Western association of physicians
Western society for clinical investigation
Plenary session
9:30 AM
Thursday, January 23, 2020

1. PREDICTIVE VALUE OF CT FOR ENDOSCOPICALLY REFRACTORY UNCONTROLLED VARICEAL BLEEDING IN CIRRHOTIC PATIENTS REQUIRING PORTAL VENOUS INTERVENTION

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10.1136/jim-2019-WMRC.1

Purpose of study: The objective of this study was to identify features on abdominal CT imaging associated endoscopically refractory uncontrolled variceal hemorrhage requiring portal venous intervention.

Methods used: Between January 2009 to January 2018, CT scans from 64 patients who experienced endoscopically refractory variceal hemorrhage requiring intervention such as transjugular intrahepatic portosystemic shunt (TIPS) placement or retrograde transvenous obliteration (RTO) and 67 patients without variceal hemorrhage but with severe symptomatic pressure gradient proven portal hypertension were analyzed. Two blinded radiologists reviewed the scans for the following parameters: size of varices, degree of variceal intraluminal protrusion, liver and splenic volumes, and portal vein diameter.

Summary of results: Gastric intraluminal variceal protrusion was found to be a strong CT parameter associated with refractory variceal hemorrhage (0.75 mm vs -2.91 mm, p=0.001). Regarding size, the hemorrhage group had larger gastric varices compared to the control group (8.03 mm vs 6.51 mm, respectively, p=0.001). However, this trend was not demonstrated in the sizes of the esophageal varices (6.28 mm vs 6.43 mm, p=0.370). Larger splenic volume (1312 cc vs 1152 cc, p=0.029) and liver volume (1514 cc vs 1143 cc, p=0.004) were also found to be predictors of endoscopically refractory variceal hemorrhage.

Conclusions: Imaging parameters on abdominal CT, such as intraluminal protrusion of gastric varices, gastric variceal size, larger splenic and liver volumes were predictive of portal venous intervention while esophageal variceal size was not.

2. RACIAL/ETHNIC DIFFERENCES IN PRENATAL AND POSTNATAL COUNSELING ABOUT MATERNAL AND INFANT HEALTH PROMOTING PRACTICES AMONG ADOLESCENT MOTHERS

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10.1136/jim-2019-WMRC.2

Purpose of study: Racial/ethnic disparities in teen birth rates persist. In 2017, the teen birth rate was 27.5 per 1000 for non-Hispanic black (NHB) teens and 28.9 per 1000 for Hispanic teens, compared with 13.2 per 1000 for non-Hispanic white (NHW) teens. Despite this difference, data describing racial/ethnic differences in counseling is lacking. This study evaluates racial/ethnic differences in 1) prenatal counseling and 2) postnatal counseling of teen mothers.

Methods used: Pregnancy Risk Assessment Monitoring System (PRAMS) 2012–2016 data, a national perinatal surveillance system was analyzed. Prenatal counseling topics included substance use (tobacco, alcohol and illicit drug use), medical care (HIV testing, influenza vaccination and recommended weight gain), and breastfeeding. Postnatal counseling topics included infant safe sleep, postpartum depression, and contraception. Bivariate associations of maternal/infant characteristics and receipt of counseling were estimated using chi-square tests. Multivariable logistic regression was used to assess the independent relationship between race/ethnicity and receipt of pre-/postnatal counseling, controlling for maternal and infant demographic and clinical factors.

Summary of results: 11805 teen mothers surveyed represent a weighted sample of 544,930, including 20.4% NHB, 29.5% Hispanic and 42.1% NHW teens. Compared to NHW teens, NHB teens were more likely to receive any prenatal substance use [AOR 8.42, 95%CI (3.47, 20.42)] and breastfeeding counseling [AOR 2.51, 95%CI (1.12, 5.65)]. Compared to NHW teens, Hispanic teens were more likely to receive any prenatal substance use (tobacco, alcohol and illicit drug use), medical care (HIV testing, influenza vaccination and recommended weight gain), and breastfeeding. Postnatal counseling topics included infant safe sleep, postpartum depression, and contraception. Bivariate associations of maternal/infant characteristics and receipt of counseling were estimated using chi-square tests. Multivariable logistic regression was used to assess the independent relationship between race/ethnicity and receipt of pre-/postnatal counseling, controlling for maternal and infant demographic and clinical factors.

Conclusions: This study demonstrates racial/ethnic differences in substance use and postpartum family planning counseling with minority teens being more likely to receive this counseling compared to their NHW counterparts. Ongoing investigation is needed to better understand the drivers of these differences including additional factors that vary by region.
EVALUATING THE PREDICTIVE VALIDITY OF THE MYHEARTSMAP PSYCHOSOCIAL SELF-ASSESSMENT TOOL IN YOUTH PRESENTING TO THE EMERGENCY DEPARTMENT WITH MENTAL HEALTH CONCERNS

10.1136/jim-2019-WMRC.3

Purpose of study The number of youth visiting the pediatric emergency department (PED) for mental health care is increasing. The use of digital self-assessment tools is an efficient method of evaluating mental health due to ease of administration without a clinician. We quantified the predictive validity of a psychosocial self-assessment tool, MyHEARTSMAP, in youth presenting to the PED with mental health concerns.

Methods used We conducted a prospective cohort study in two tertiary PEDs from December 2017-June 2019. Youth 10–17 years old triaged for a mental health concern were screened and enrolled to complete MyHEARTSMAP on a mobile device. A blinded clinician conducted a HEARTSMAP assessment, which was the reference standard. Both tools collect information about ten sections: Home, Education and activities, Alcohol and drugs, Relationships and bullying, Thoughts and anxiety, Safety, Sexual health, Mood and behavior, Abuse, and Professional resources. Based on scoring from 0–3 (no concern, mild, moderate, or severe), a score for psychosocial domains (psychiatry, social, youth health, and function) is computed and resources are recommended based on need. Sensitivity and specificity were measured for youth and guardian MyHEARTSMAP assessments in detecting severe psychiatric concerns and any other domain concerns.

Summary of results We screened 510 youth and 167 families enrolled. Concerns included suicidality or safety concerns (49.7%) and mood (25.7%) and behavioral disorders (12.6%). The remaining were substance use, social issues, and eating and thought disorders. Youth and guardian’s sensitivity for detecting a severe psychiatric concern was 70.5% (95% CI: 54.8%, 83.2%) and 80% (65.4%, 90.4%) respectively. Specificity was 34% (24.6%, 44.5%) and 41.2% (31.5%, 51.4%). The sensitivity for detecting any domain concern in youth and guardians was 93.9% (87.3%, 97.7%) and 93.3% (86.8%, 97.3%) for social, 88.2% (80.4%, 93.8%) and 90% (82.8%, 94.9%) for youth health, and 96.8% (91.9%, 99.1%) and 97% (92.4%, 99.2%) for function.

Conclusions Assessment by families using MyHEARTSMAP is sensitive for detecting psychosocial concerns requiring mental health follow-up beyond PED evaluation.

INTEGRATING TECHNOLOGY INTO PATIENT EDUCATION TO EMPOWER PEDIATRIC ENDOCRINOLGY PATIENTS TO IMPROVE THEIR HEALTH THROUGH A PATIENT-CENTERED CARE APPROACH

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Purpose of study Dybiosis of the human microbiome, caused especially by a decline in its microorganism diversity, can play a role in the development of harmful, long-term conditions that impact human health. A major contributor to this dysbiosis is the use of antibiotics to treat bacterial infections. Antibiotics can be divided into two groups: narrow spectrum antibiotics, which are effective against a specific group of bacteria, and broad spectrum antibiotics (BSA), which are effective over a wider range of bacteria and can pose a risk of disrupting the microbiome. Antibiotic treatment of Group A strep (GAS) infection, which causes strep throat and scarlet fever, serves as area where microbiome harm can be reduced via treatment with penicillin (PCN), a narrow spectrum drug that GAS are 100% sensitive to. Antibiotics commonly used for treatment, however, include amoxicillin, a moderate BSA, along with other broader spectrum antibiotics. We sought to characterize the antibiotics used to treat strep throat and scarlet fever and to further describe the extent to which BSA are used for treatment.

Methods used We performed retrospective data extraction and analyses of 3,169 patients who were less than 22 years old, received a diagnosis of strep throat or scarlet fever, and were seen in an outpatient setting. Patients were further excluded if they had a PCN allergy or had any diagnoses of co-infections or comorbidities that would require the use of BSA. We then used descriptive statistics as our primary mode of data analysis.

Summary of results In summary, 2,208 patients were determined to have simple strep throat or scarlet fever. Of these patients, 65% were treated with various BSA, mainly amoxicillin; 27% were treated with PCN; and 8% were given no antibiotics. With subjects divided by age group (0–3, 4–12, 13+), the trend of BSA being used more often for treatment continues, though a greater proportion of patients aged 13 and above were found to be treated with PCN.

Conclusions Our findings highlight opportunities for clinicians to reflect on their use of antibiotics in treating strep throat and scarlet fever. It is important for providers to prioritize the reduction of harm to the microbiome, which is essential in maintaining human health, by reducing their use of BSA whenever possible.
providing information in a multimedia format, and as such can serve as a supplement to the more traditional forms of education in order to improve self-efficacy in patients and their families.

**Methods used** Two high school students enrolled in the UCSF Fresno Medical Education Program Summer Biomedical Research Internship Program were charged with the creation of the websites. 34 diabetes patients and 47 patients with other endocrine disorders were surveyed, along with their parents. Websites were built using Google Sites given its easy accessibility, with all content either original or with Creative Commons licensing. Videos were created using iMovie and housed in a YouTube site.

**Summary of results** 77% of diabetes patients/parents and 64% of other endocrine patients/parents were interested in the development of educational websites. Two separate websites were created: one for diabetes and for patients with other endocrine disorders. Both websites contain easy-to-understand information written for those even with lower literacy levels. Brief educational videos were created to reinforce concepts taught in the clinic, and diabetes videos such as ‘My Diabetes Monster’ were created by our patients and parents to empower them and show them that they are not alone in their struggles with diabetes.

**Conclusions** The majority of patients and parents surveyed expressed interest in the development of easily accessible educational websites, which can be easily accessed by both inpatients and outpatients even outside normal business hours. Such sites can be built by high school students interested in health careers under the guidance of the physician, thus providing medical pipelines as well as patient education. These sites should help patients and their families improve their self-efficacy in providing care for their disease.

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**6 THE EFFECTIVENESS OF AN ADOLESCENT LONGITUDINAL EVALUATION OF PEDIATRIC RESIDENCY DIDACTICS TRANSITION FROM NOON CONFERENCE TO AN ACADEMIC HALF DAY**

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**Purpose of study** In California’s Central Valley, the teen birth rate is almost double the national average and sexually transmitted infection (STI) rates are disproportionately higher amongst minority populations. In this study we assessed the effectiveness of an adolescent reproductive health symposium for interprofessionals in offering comprehensive reproductive services to adolescents in this region.

**Methods used** The sample consisted of interprofessionals, including pediatricians, pediatric residents, nurses, and community healthcare educators. Participants were recruited from a one-day, lecture style symposium where education was provided on national and local current trends in STI and pregnancy rates amongst adolescents, effective contraceptive methods, community resources, and referral options to provide comprehensive reproductive healthcare services to adolescents. A pre and post survey instrument and a 3-month post survey were utilized to assess provider perceptions of providing comprehensive care for adolescents.

**Summary of results** Participant perceptions of their clinical competencies in offering comprehensive adolescent reproductive health services improved after the symposium (n=64). Statistically significant differences were noted in the perceptions of having adequate training (p<0.001) and feeling comfort taking a psychosocial and sexual history (p=0.001). There were also statistically significant differences in the perceptions of being adequately prepared to counsel adolescent patients on contraceptive methods (p<0.001), including options for long acting reversible contraceptives (p=0.003) and local referral options (p<0.001). There were no statistically significant differences found 3 months after the initial education; however, participants showed maintenance of perceptions.

**Conclusions** A symposium based intervention demonstrates the positive impact that interprofessional education has on adolescent reproductive health by increasing provider competency in offering comprehensive adolescent reproductive health services.

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**7 LONGITUDINAL EVALUATION OF PEDIATRIC RESIDENCY DIDACTICS TRANSITION FROM NOON CONFERENCE TO AN ACADEMIC HALF DAY**


**Purpose of study** In 2009, the University of New Mexico (UNM) Pediatric Residency transitioned program didactics from noon conference to an academic half day. Immediate evaluation of resident and faculty satisfaction, resident attendance, board exam pass rate and In-Training-Exam (ITE) scores showed improvements with this change. These results by Dr. Walter Dehority were published in 2012. The primary aim of this study was to investigate the longitudinal impact of the academic half day at UNM on resident and faculty satisfaction, resident attendance and clinical knowledge in the form of standardized exam results. In addition, use self-determination theory to explore effects of the academic half day on autonomy, competence and relatedness in pediatric residents.

**Methods used** In this mixed methods study, surveys regarding satisfaction were conducted of current pediatric residents (n=32) and faculty (n=32) at UNM. To assess clinical knowledge, quantitative measures such as pediatric board exam pass rate were evaluated. Finally, focus groups were conducted with pediatric residents and faculty separately to evaluate concepts related to resident autonomy, competence and engagement with the academic half day.

**Summary of results** Resident and faculty satisfaction was sustained with significance at 88% and 67% (p-value of 0.02 and 0.009), respectively. Resident attendance was also sustained at 88.4% (p=0.005). Board exam pass rates are increased since 2009, with unclear significance. Focus groups...
Identify overall satisfaction from residents and faculty and identify a need for addressing resident engagement with didactics secondary to clinical pressures and interactivity of didactics.

Conclusions The academic half day shows continued success in terms of faculty and resident satisfaction and resident attendance. More research is needed to investigate other factors related to the board pass rate. Further intervention and research will be needed to improve resident engagement in the academic half day.

Purpose of study Current curriculums teach medical trainees how to talk with patients and families though online videos, simulated patients, small groups, and clinical encounters, with or without feedback given by trained providers. Few pediatric curriculums address communication fundamentals: semantics, phrasing, and developmental milestones. The purpose of this curriculum is for pediatric residents to learn effective semantic and phrase usage as well as understand developmental milestones to improve their communication knowledge and skill confidence.

Methods used Pediatric residents of all training levels at one academic institution participated in a one-time 45-minute workshop. The curriculum included three exercises covering semantics, phrasing and milestones via small group discussions and self-assessments of communication/milestone knowledge. Pre-, post-, and 7-month intervention surveys that evaluated knowledge and confidence about their communication skills with pediatric patients were collected. Qualitative feedback was acquired for curriculum improvement.

Summary of results Forty-one pediatric residents participated in the workshop: 71% female, 88% categorical pediatrics, and 17% with children. Residents included 1st, 2nd, 3rd, and 4th years: 34%, 44%, 20%, 2%, respectively. Survey participation (3 surveys) averaged at 68%. Significant improvement occurred following the workshop: knowledge about speaking with patients increased from 9% to 44% (p<0.01, Z-score=-3.1), knowledge in developmental stages increased from 28% to 72% (p<0.01, Z-score=-3.5), confidence while speaking with patients increased from 16% to 53% (p<0.01, Z-score=-3.2). At 7-month follow up, 91% of participants stated that they used at least one piece of knowledge or skill gained from this workshop.

Conclusions This workshop increases communication knowledge and confidence immediately after as well as 7 months after the session by focusing on the semantics, phrasing, and developmental milestones. It provides pediatric trainees with the fundamentals for communicating with pediatric patients, which are applicable for all patients, compared to learning through experience in feedback, which may be patient-specific.

Purpose of study Distal radius and ulna buckle fractures are a frequent reason for pediatric emergency department (ED) visits. Previously, providers used immobilization in a short arm cast. Studies have shown equivalent outcomes in using a removable splint over a cast, with reduced costs, increased...
efficiency, and improved patient/family satisfaction. Our baseline data demonstrated that nearly all children with distal radius/ulna buckle fractures were being casted at our ED. We planned to increase the use of removable splints in children 2–18 years old with these fractures from 17% to >90% and to reduce orthopedic consultations in the ED from 88% to <10% over a 12 month period.

Methods used We performed a retrospective chart review to obtain baseline data. In collaboration with orthopedics, a standardized protocol was created and implemented in January 2019. Treatment involved placing splints for patients ages 2–18 with distal forearm buckle fractures seen on x-ray without consulting or following up with orthopedics. Emergency medicine, pediatric, and orthopedic physicians were educated on the protocol. Standardized instructions for caregivers and primary care providers were made. We reviewed monthly data on the number of splint placements, orthopedic consultations and follow-ups, x-rays, and length of stay.

Summary of results In the first seven months of implementation there have been 11 patients that fit the inclusion criteria. The proportion of removal splints increased from 17% (n=9) to 63% (n=7) (p=0.01). Orthopedic and plastic surgery consultations decreased from 88% (n=46) to 36% (n=4) (p=0.02), and surgical clinic follow up decreased from 86% (n=45) to 36% (n=4) (p=0.008). Average x-rays done decreased from 1.3 to 1.0 (p=0.02), and length of stay decreased from 202.9 to 154.3 minutes (p=0.0001).

Conclusions Initiation of a protocol for distal forearm buckle fractures in our pediatric ED has increased the use of removal splints and reduced orthopedic consultations, clinic follow up, number of x-rays, and length of stay. We plan to increase the duration of data collection to achieve adequate sampling. Next steps include educating plastic and trauma surgery services about the protocol and further partnering with caregivers to prevent unnecessary orthopedic referrals.

Cardiovascular I
Concurrent session
12:45 PM
Thursday, January 23, 2020

T-CELL DEFICIENT MICE DEMONSTRATE REDUCED
ATHEROSCLEROTIC PLAQUE BURDEN

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Purpose of study Atherosclerosis is characterized by arterial plaque deposition. The CD40-CD154 inflammatory dyad is a major driver of the auto-inflammation seen in atherosclerosis and type 1 diabetes (T1D). A sub-type of CD4+ T cells that express CD40 (Th40 cells) is increased in T1D, but has not been defined in atherosclerosis. Thus, this study seeks to explore the role of this pathogenic T-cell on plaque formation.

Methods used ApoE−/− transgenic mice represent a model of atherosclerotic disease. We have generated a novel TCRαβ−/−
Apoe-/- mouse model to investigate the role of Th40 cells. Baseline and ‘add back’ T cell experiments have been conducted. Mice are sacrificed at 8 months of age and then dissected to obtain the aortas and hearts. The aorta is used for en-face Sirius Red stain analysis while serial aortic valve cross sections are used to characterize the lesion in terms of area, volume, necrosis, and cellular content.

**Summary of results** Trichrome-stained aortic valve cross sections show a significant reduction in overall plaque as well as ability to form advanced plaque (as determined by macrophage content) in the Apoe-/- TCRd/-/- mouse model (shown in image B) as compared to its Apoe-/- counterpart (A). This reduction in plaque deposition is also shown in the Sirius Red analysis of the whole aorta. Representative samples of double KO mice show 16% plaque deposition (C) compared to 33% within Apoe-/- mice (D).

**Conclusions** T-cell deficient Apoe-/- mouse models demonstrate a decreased plaque burden in comparison to the Apoe-/- counterparts illustrating that T-cells are implicated in atherosclerosis pathogenesis. Future studies will characterize the cell populations within the plaques.

### 13 WARFARIN ANTIMOULATION FOLLOWING PEDIATRIC HEART SURGERY: COMPARING HOME VERSUS TRADITIONAL LAB MONITORING

| A Lippay* | D Griffin, *J Frazier, L Sah. University of California San Diego, San Diego, CA; 2Rady Children’s Hospital, San Diego, CA; 3University of California Los Angeles, Los Angeles, CA; 4Santa Barbara Cottage Hospital, Santa Barbara, CA |

**Purpose of study** Warfarin is commonly used following placement of mechanical heart valves or an extracardiac conduit in the Fontan procedure in order to minimize risk of thrombotic events. The degree of anticoagulation is monitored by the patient’s international normalized ratio (INR), which is traditionally done by serial outpatient blood draws. The goal of this pilot study is to determine if anticoagulation can be better managed using a home INR monitoring system compared to traditional laboratory monitoring.

**Methods used** Patients who underwent artificial mechanical valve placement and/or an extracardiac Fontan procedure were consented and randomized to one of two 3-month arms: INR testing via 1) traditional laboratory monitoring (venipuncture), or 2) a home INR monitor (finger stick test). After 3 months, patients were crossed-over to the other arm (6 months total in the study). Frequency of INR test was determined per cardiologists’ clinical judgement. A questionnaire was completed after each arm of the study. Eleven subjects have been enrolled into the study thus far: 10 have completed at least one arm of the study, 5 have completed both arms, and 1 patient was withdrawn after the first arm due to non-compliance. Student t-tests were performed to compare time and cost differences between the two arms.

**Summary of results** The ease of obtaining INR value between home and laboratory monitoring was rated as similar (3.4/5 vs 3.8/5), with similar pain rating experience by the subject (2.5 vs 2.4/5). However, parents were more satisfied with using the home INR system (4/5 vs 2.9/5) and were more likely to recommend to others (4.2/5 vs 2.5/5). The average time it took to use the home INR monitor versus lab monitoring was also significant (3.8 minutes vs 53.3 minutes; p<0.0001). The percentage of INR in therapeutic range for subjects was also similar between arms (home 46.9% vs lab 43.4%; p=0.6396).

**Conclusions** Overall, preliminary data show similar outcomes between home vs lab INR monitoring, with the former showing improved patient satisfaction and time saved. Ongoing cost-effective analysis is being conducted as the pilot study continues.

### 14 ACTIVE SMOKING IS ASSOCIATED WITH DECREASED RATES OF WOUND HEALING AFTER ENDOVASCULAR TREATMENT OF CRITICAL LIMB ISCHEMIA

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10.1136/jim-2019-WMRC.14

**Purpose of study** To examine whether active smoking is associated with worse complete wound healing rates in patients with CLI undergoing endovascular interventions.

**Methods used** A single center observational study was conducted. Consecutive patients with CLI who underwent endovascular interventions were included. Smoking status was assessed at the time of the intervention and active smoking was compared to no active smoking at the time of the intervention. Cox and logistic regression analyses were conducted to compare the complete wound healing rates of the two groups during follow-up.

**Summary of results** In total, 264 patients (active smokers: n=41; not active smokers: n=223) and 553 lesions were included. Active smoking was associated with decreased rates of complete wound healing in the 6-month univariate cox regression analysis (HR: 0.22; 95% CI: 0.07 - 0.71; p=0.012). The 6-month KM estimates for complete wound healing were 8.9% for the active smoking group vs 34% for the no current smoking group. Active smoking was also associated with decreased rates of complete wound healing in the 9-month univariate cox regression analysis (HR: 0.43; 95% CI: 0.21- 0.90; p=0.026) and multivariate analysis (HR: 0.11; 95% CI: 0.01 – 0.94; p=0.044). The 9-month KM estimates for complete wound healing were 25.2% in the active smoking group vs 46% in the no active smoking group. No difference was found between the two groups in the 9-month major amputation rates. The time to complete wound healing was longer in the active smoking group (269.7±187.5 days vs. 208.3±115.7 days; p=0.009).

**Conclusions** Active smoking status at the time of intervention in CLI patients is associated with lower rates of complete wound healing during 6- and 9-month follow-up. Active smoking was associated with a two-month delay in the median time to wound healing. These findings emphasize the importance of smoking cessation for patients with CLI.
Walter: Hospital Reporting of Critical Congenital Heart Disease Screening

California passed legislation in 2013 mandating oxygen saturation (SpO2)-based screening for critical congenital heart disease (CHD) screening and annual data submission to the Department of Health Care Services (DHCS). We conducted a retrospective review of CHD screening data reported to California DHCS to describe reporting compliance and assess data accuracy.

**Methods used**
This was a retrospective review of hospital CHD screening reports provided to California DHCS from January 2015 to December 2016. We compared the reported birth data to California’s Office of Statewide Health Planning and Development (OSHPD) for accuracy of live births reported and to determine frequency of reporting from hospitals meeting the CHD screening reporting requirements. We also evaluated the data for other potential markers of accuracy using proxies such as number of report revisions required and accurate summation of variable cells (i.e. all screen results accounted for, or all eligible newborns undergoing screening).

**Summary of results**
Thirty-four percent of hospitals did not comply to mandated CHD screen reporting. When hospitals comply to reporting, 45.8% in 2015 and 44.2% in 2016 submitted reports in which the number of completed screens did not match the number of screen results. Only 21.8% in 2015 and 21.1% in 2016 of hospitals submitted live births that matched their respective live births reported to a second database. In 2015, the submitted data represented 375,283 live births, which was 80% of the 468,936 live births reported to OSHPD that year. In 2016, the submitted data represented 318,424 live births, or 56% of the 570,786 live births reported to OSHPD that year. For the two years combined, up to 623 newborns with CHD could have been missed or unaccounted for due to non-reporting.

**Conclusions**
A third of hospitals do not comply to reporting guidelines and over half of the submitted reports are inaccurate. A large number of newborns with CHD may potentially be missed due to under/inaccurate reporting.

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Vlad: Variability of End-Tidal Carbon Dioxide in Out-of-Hospital Cardiac Arrest

We hypothesized that molecular mechanisms of HF pathogenesis differ across the life-course, a hypothesis which we tested with a mouse model of cardiac dysfunction at three distinct stages of life.

**Methods used**
C57BL/6 mice at pediatric (5 weeks; n=12), adult (3–5 months; n=12), and old (18 months, n=10) ages were treated with a subcutaneous mini-osmotic pump that eluted isoproterenol (ISO; 30 mg/kg/hour), a non-selective β-adrenergic receptor agonist commonly used to induce acute cardiomyopathy in mice. Following 6 days, we performed echocardiographic, biochemical assessments, and RNA sequencing of the left ventricle (LV).

**Summary of results**
Both the pediatric and adult groups underwent hyperthermic remodeling in response to ISO, as evident by higher LV weight relative to tibia length (TL). However, ISO exposure did not increase LV/TL in old mice. Echocardiographic imaging demonstrated thickening of the ventricular wall in ISO mice compared to control. Expression of pro-fibrotic mediators also differed across the life-course in response to ISO, with adults inducing a pro-fibrotic transcriptional program (α-smooth muscle actin, fibronectin, collagen, peristin) that was attenuated in old and absent in pediatric animals. RNA-sequencing identified that 119, 1515, and 33 genes were significantly differentially expressed in pediatric, adult, and old mice exposed to ISO, respectively (p<0.05). Of these genes, only 2 transcripts were differentially expressed across all three ages. Therefore, biological age significantly impacts the molecular mechanisms of ISO-induced cardiac remodeling. Ongoing analysis of these molecular targets will inform HF therapies using age as a biological variable.
KENCOR REMOTE MONITORING SYSTEM OPTIMIZES BLOOD PRESSURE CONTROL AND IMPROVES SERUM CREATININE LEVELS IN HYPERTENSIVE PATIENTS

1K Dhond*, 2Y Gill, 2M Villalon, 3S Lahsaei, 2MJ Dhond, 2C Mancherje, 2J Breneisen, 2M Dhond, 1UC Santa Cruz, Santa Cruz, CA; 2Northbay Healthcare, Fairfield, CA and 2UC Davis, Davis, CA

Purpose of study Hypertension (HTN) costs the US $48.6 billion per year, it occurs in one third of the US population with only 54% reaching guideline values. We used KenCor Remote Monitoring System (KRMS) to evaluate its utility in optimizing blood pressure control.

Methods used 43 clinic patients with HTN were enrolled in KRMS. Patients were oriented into KRMS software which was downloaded onto their smartphones. Patients were provided with a Bluetooth-enabled blood pressure (BP) cuff and weigh scale. Patients completed a daily questionnaire and recorded their vitals, with uploading of data to the clinic nurse. KRMS then stratified the patients into low (green), medium (yellow), or high (red) risk vitals. Patients were compliant with KRMS.

Summary of results Of 353 included patients, initial ETCO2 were 24 mmHg [11/34] and 10 mmHg [2/22] for shockable and non-shockable rhythms, respectively. Pre-intubation values for these cohorts were 27 mmHg [4/40] and 10 mmHg [3/27] and post-intubation values were 43 mmHg [33/59] and 46 mmHg [25/64]. Post-defibrillation, ETCO2 increased from 38 mmHg [30/53] to 54 mmHg [41/60] among patients who achieved ROSC and 41 mmHg [25/57] to 42 mmHg [27/56] among those without ROSC. With initial VF/VT, higher post-intubation and post-ROSC ETCO2 were associated with worse cerebral performance categorization (CPC) among survivors (post-intubation CPC 1/2: 35 mmHg [28/44], CPC 3/4: 59 mmHg [48/71]; post-ROSC CPC 1/2: 50 mmHg [45/59], CPC 3/4: 64 mmHg [57/67]).

Conclusions Patients with ROSC after defibrillation had larger rises in ETCO2 than patients without ROSC. Pre-intubation ETCO2 values appear clinically unreliable. Our finding that higher ETCO2 is associated with worse neurologic outcomes for patients who survive to hospital discharge warrants further investigation.

NSTEMI AND HREF DUE TO METHAMPHETAMINE

JA Coleman*, D Aguirre. Kern Medical, Bakersfield, CA

Purpose of study To discuss the diagnosis, evaluation, and management of methamphetamine-induced acute myocardial infarction in the presence of methamphetamine induced cardiomyopathy.

Methods used Retrospective Chart Review.

Summary of results A 35-year-old male with no pmh who has a social history significant for tobacco, alcohol, methamphetamine, cocaine, and PCP abuse presented to the emergency department 3 days h/o dyspnea and acute onset of new onset of pressure-like pain in his chest, which was constant, 7/10, and non-radiating, wth no alleviating or aggravating factors. Upon presentation he was hemodynamically stable, and admitted to using methamphetamine hours prior. Chest x-ray was unremarkable. EKG showing no ST segment elevations or depressions. Initial serum troponin was elevated at 0.63 and urine toxicology was positive for methamphetamine and PCP. NSTEMI was diagnosed. He was admitted and started on ACS protocol, and TTE ordered. Repeat troponin was 2.45, and TTE revealed a LVEF of 25–30%. Patient subsequently underwent a left heart catheterization which found the proximal left anterior descending (LAD) artery to have a lesion of 80% with the left ventriculography estimating a LV EF of 20%. Patient was transferred to MICU while pending transfer to tertiary institution for PCI. Patient underwent revascularization and bare metal stenting of proximal LAD baring no complications and closely followed up in cardiology clinic.

Medications on discharge were aspirin, clopidogrel, carvedilol, lisinopril, furosemide, atorvastatin, and spironolactone. Eight months later, the repeated TTE resulted in significant improvement of the LV EF of 40–45% from 20–25%. The dual antiplatelet therapy with aspirin and clopidogrel was given for a duration of 12 months. Patient is now 14 months post PCI and is remaining clinically asymptomatic.
Conclusions Amphetamine associated acute myocardial infarction may become more common if the rate of amphetamine abuse continues to increase. Methamphetamine associated cardiomyopathy appears to be potentially reversible upon cessation of methamphetamine and guideline directed management for heart failure.

Endocrinology and metabolism I
Concurrent session
12:45 PM
Thursday, January 23, 2020

20 UNMASKING OF PNEUMOCYSTIS JIROVECI PNEUMONIA TRIGGERED BY TREATMENT OF CUSHING’S SYNDROME
SC Fernandes, C Schmid*, AM Mansoor, E Varlamov. Oregon Health and Science University, Portland, OR
10.1136/jim-2019-WMRC.20

Case report Case: A previously healthy 29 year old man presented with new onset hypertension, diabetes mellitus (DM), easy bruising, 70-lbs weight gain and painful purple stretch marks. Physical examination was notable for blood pressure of 156/91 mmHg, moon facies, facial plethora, acne, dorso-cervical and supraclavicular fat pads, violaceous striae, abdominal bruising and thin skin. Labs showed hypokalemia (3.1 mmol/L), markedly elevated adrenocorticotropic hormone (ACTH 480 pg/mL; Nl <45) and 24 hour urine free cortisol (UFC 7950 ug/day; Nl <60), consistent with an ACTH-dependent Cushing’s syndrome (CS). Pituitary MRI showed a possible 4 mm lesion, however inferior petrosal sinus sampling confirmed ectopic ACTH CS. CT and nuclear imaging were negative. Escalating doses of ketoconazole were started with UFC reduction to 73.3 ug/day. He developed transaminitis, as well as respiratory distress with hypoxemia. Chest imaging demonstrated bilateral ground glass opacities concerning for Pneumocystis jiroveci pneumonia (PJP). Serum 1,3 beta-D glucan was elevated (188 pg/mL; NL <80) increasing suspicion for PJP. Patient declined sinus sampling to confirm diagnosis and was empirically treated with trimethoprim-sulfamethoxazole and prednisone, resulting in rapid improvement of respiratory symptoms. The patient underwent bilateral adrenalectomy for CS and started on replacement hydrocortisone and fludrocortisone. He had 30 pound weight loss and diabetes and hypertension resolved in 3 weeks.

Conclusions Patients with CS are at high risk for opportunistic infections. Cortisol excess depresses immune function allowing PJP to colonize the lungs. Treatment with cortisol decrease results in T cells recovery, which can cause an inflammatory reaction to PJP, analogous to immune reconstitution syndrome in HIV patients. Ectopic CS and higher cortisol levels increase PJP risk. Thus, PJP prophylaxis should be considered in patients with severe CS prior to initiation of any cortisol lowering therapies.

21 EVALUATION OF THE IMMUNE SYSTEM IN PEDIATRIC CUSHING DISEASE
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10.1136/jim-2019-WMRC.21

Purpose of study Cushing Disease (CD) is a rare pediatric endocrine disorder that results from adrenocorticotropic hormone (ACTH) secreting pituitary adenomas. CD should be recognized as early as possible in the pediatric population due to the significant morbidity and mortality associated with late diagnosis and treatment. In the present study, we sought to describe the relationship between complete blood count (CBC) values, serum cortisol level, and size of the thymus. We also studied the association of the neutrophil to lymphocyte ratio (NLR), recently identified as a diagnostic and prognostic factor in cancer, with markers of hypercortisolism.

Methods used The inclusion criteria for the study included diagnosis of CD before age 18 years, diagnosis between 1997 and 2019, and previous imaging study that included visualization of the thymus (obtained prior to transphenoideal surgery). CBC and serum cortisol levels at diagnosis were retrieved from medical records via retrospective chart review. Neutrophil and lymphocyte z-scores were calculated based on age and sex specific reference ranges. NLR was calculated using absolute levels among patients 6–12yo and >12yo. All studies were approved by the NIH Institutional Review Board.

Summary of results Among 48 patients that met our inclusion criteria, no significant correlation was found between Hemoglobin (Hb) and cortisol. A significant correlation was found between cortisol levels and neutrophil (β=0.07, R2=0.1978, p<0.001) and lymphocyte (β=0.04, R2=0.1524, p<0.01) z-scores. Midnight cortisol was significantly correlated with the NLR in patients >12 years old (β=0.14, R2=0.6958, p<0.0001). Results of NLR for patients ≤12 years old did not reach statistical significance (p=0.096). Thymic data are pending complete analysis.

Conclusions Immune dysregulation in CD can be used as supportive evidence of its diagnosis. NLR can be easily calculated from CBC and correlates with cortisol with a stronger coefficient of determination than solely using neutrophil or lymphocyte values. Further investigation is needed to establish a cutoff for NLR that can potentially differentiate patients with CD from eucortisolemic obese patients.

22 TESTOSTERONE AND CORTISOL MODULATE THE METABOLIC EFFECTS OF SLEEP RESTRICTION
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10.1136/jim-2019-WMRC.22

Purpose of study Sleep restriction is highly prevalent and increases the risk of type 2 diabetes mellitus. The mechanisms by which sleep restriction causes insulin resistance (IR)
are not known. Understanding these mechanisms would allow targeted interventions beyond requiring more sleep. Sleep restriction decreases testosterone, and increases cortisol, which are the main anabolic and catabolic hormones, respectively. Through a dual hormonal clamp intervention, we investigated whether testosterone and cortisol affect IR due to sleep restriction.

Methods used 34 healthy men aged 33.3±6.4y with BMI of 25.4±2.5 kg/m² completed the study. Subjects were admitted to the Clinical and Translational Research Center. They had one night with a 10h opportunity (22:00–08:00) for baseline sleep and were then restricted to 4h sleep (01:00–05:00) for 4 consecutive nights. This pattern was repeated on two separate occasions with different treatment conditions, in randomized order: dual testosterone/cortisol clamp (ketoconazole + transdermal testosterone gel + oral hydrocortisone at mid-physiological doses) during all sleep restriction days, or no clamp (matching placebo). A 3-hour intensively sampled (23 samples) oral glucose tolerance test was performed after the baseline night, and after the fourth sleep-restricted night, to calculate IR by Matsuda Index (Mi) and by minimal model (Si). Effects of condition and measurement day and their interaction were determined by mixed-model.

Summary of results Following sleep restriction, Mi showed greater IR in both conditions, but the increase was dampened in the dual clamp condition (F=4.90, p=0.029 for interaction). A composite sensitivity index that combined all glucose metabolic indices (Si, HOMA-IR, Matsuda Inex) confirmed this finding (F=4.81, p=0.029 for interaction). Conclusions A dual hormonal clamp intervention of testosterone and cortisol in a sample of healthy, young adult men dampened the negative effects of sleep restriction on glucose metabolism. Our results show that testosterone and cortisol modulate the effects of sustained sleep restriction on insulin resistance.

Methods used Two randomized, double-blind, placebo-controlled studies in healthy men were previously performed to assess the safety and tolerability of DMAU and 11β-MNTDC taken orally for 28 days. Insulin and adiponectin assays were performed on banked samples. Changes in weight, LDL-C, HDL-C, fasting glucose, HOMA-IR, and adiponectin were assessed. Log transformations were performed as needed. Two-way ANOVA with post hoc Tukey HSD was performed to assess for dosage (0, 200, or 400 mg) and drug (DMAU or 11β-MNTDC) effects.

Summary of results A total of 85 subjects were included in this analysis. Overall, there was a significant decrease in HDL-C and increase in weight and LDL-C (all p<0.001). For weight and HDL-C, both the 200 mg and 400 mg groups were significantly different from placebo, but there was no significant difference between the 200 mg and 400 mg groups. For LDL-C, the 200 mg groups were not significantly different from placebo (p=0.059) but the 400 mg groups were significantly different from placebo (p<0.001); there was no significant difference between the 200 mg and 400 mg groups (p=0.11) nor differences between the two novel androgens. There was no difference in fasting glucose, adiponectin or HOMA-IR.

Conclusions DMAU and 11β-MNTDC reduced HDL-C and increased weight and LDL-C in these analyses. Changes in metabolic parameters may be considered during further development of male hormonal contraception.

Purpose of study Pheochromocytoma is a rare tumor originating in chromaffin cells of the adrenal gland with an incidence rate estimated at 0.8 per 100,000 individuals per year. There have been few documented cases of the tumor and its consequences when not recognized in time.

Methods used Retrospective case report.

Summary of results A 45-year-old male with unknown past medical history presented via emergency medical services (EMS) to the emergency department (ED) after patient was found down by the railroad with reported right facial droop, dysarthria, and weakness. EMS reported that a friend on scene disclosed that patient had recently smoked methamphetamine. On arrival to the ED, patient was noted to have a blood pressure of 265/131, with a National Institutes of Health Stroke Scale (NIHSS) of 10 and a Glasgow Coma Scale (GCS) 14. He was started on intravenous blood pressure medications and later switched to oral regimen. CT brain found hypodensities in basal ganglia, periventricular centrum semiovale, regions worse on left, suggestive of lacunar infarctions. Hospital course was complicated by uncontrolled hypertension, right sided paresis, aphasia and dysphagia. Given high risk for aspiration, the decision was made to place a percutaneous gastrostomy tube for nutrition and medication administration and started on warfarin for its presumed atrial fibrillation.
management. CT abdomen and pelvis performed prior to procedure revealed an incidental finding of a 6 cm heterogeneous adrenal mass likely pheochromocytoma confirmed by serum metanephrine assays greater than 1000 pg/mL. The patient was treated with appropriate antihypertensive medication based on guidelines set forth by Endocrine Society, with planned laparoscopic resection of adrenal mass scheduled for two weeks later.

Conclusions Given how rare the diagnosis of pheochromocytoma is, the differences in presentation and the need for early diagnosis and management, a clinician’s index of suspicion should be higher in individuals with no risk factors presenting with neurological manifestations and uncontrolled hypertension.

Case report A 15 year old hispanic male was in his usual state of health until morning of presentation. He awoke to go to the bathroom, but found himself unable to move his extremities well. He then proceeded to crawl to the bathroom where he vomited twelve times. In ED, vitals were normal, but exam remarkable for flaccid weakness of upper and lower extremities, and mild proptosis. Pertinent labs revealed potassium 1.4, glucose 169, magnesium 1.6, pH 7.41. EKG illustrated prolonged QT interval, ST depression and U waves. He was given IV potassium over four hours with repeat potassium 4.8. Repeat EKG normalized with paralysis slightly improved. Toxicology initially consulted for possible ingestion cause, but no substance found in urine or blood. Thyroid studies revealed TSH of <0.01, and elevated free T4 of 3.83, confirming the diagnosis of thyrotoxic periodic paralysis (TPP). Propranolol was started, along with supplemental magnesium and potassium. Within 24 hours of presentation, patient’s paralysis completely resolved. Patient was found to have Graves’ disease with elevated TSI at 133, and diffuse uptake in radionuclide thyroid scan. He was subsequently started on methimazole. The pathophysiology of TPP is unclear but is theorized to rapid intracellular shifts of potassium, or catecholamine release from the vomiting.

Conclusion The patient in this case demonstrates a departure from the usual cases of TPP. He was a Hispanic male, whereas there is normally a higher incidence amongst Asians. The patient was also 15 years old, which is younger than the cited peak ages of 20–40. Sometimes there can be a genetic predisposition, as paternal grandmother possibly had the same symptoms. More often, these children have no preceding disease, as in this case. It is important for ED providers and pediatricians alike to keep this in mind as it can be easily remedied, but fatal if not recognized.
Abstracts

Healthcare delivery research I
Concurrent session
12:45 PM
Thursday, January 23, 2020

27 LIGHTS, CAMERA, SURGERY: VIDEO RESOURCES ENHANCING HEALTHCARE COMMUNICATION
K Nguyen*, C Binda, P Mangat, D Duffy, K Mulpuri. University of British Columbia, Vancouver, BC, Canada; BC Children’s Hospital, Vancouver, BC, Canada; University of British Columbia, Vancouver, BC, Canada

Purpose of study Effectively communicating medical information to patients and their caregivers can help improve patient understanding, adherence to treatment and health outcomes. Despite the importance of communication, physicians often do not communicate satisfactorily with their patients. Our purpose is to create interdisciplinary educational video resources for caregivers who are accessing surgical services at BC Children’s Hospital (BCCH). These videos, supplementing regular clinical practice, will help patients and caregivers understand their child’s condition, improve surgical outcomes, reduce stress as well as the financial burden of unnecessary emergency room visits.

Methods used Surgical team members and research staff from three departments, orthopaedics, urology and surgery at BCCH, identified suitable video topics and learning objectives. Existing patient resources were combined with further research and expert opinion to draft video scripts. Patients and caregivers were voluntarily asked for their participation and consented through signing a media release form. Videos were then created and edited using software such as Powtoons, Wire-Wax and Adobe After Effects. Completed videos were made available on the Office of Paediatric Surgery Evaluation and Innovation (OPSEI) YouTube channel and on clinic websites. URLs and QR codes were also used to connect viewers to the video resources.

Summary of results 62 videos were created in collaboration with 31 families, 7 surgeons, 3 physiotherapists, 2 occupational therapists, 5 nurses, and 5 researchers. Video topics for each series included hip dysplasia, clubfoot, limb reconstructions, cystectomy, ileus from nephrectomy cases, along with reductions in complication rates and ileus for nephrectomy cases over the year of the consultation service. The primary outcome marker was a reduction in length of stay. Secondary outcome markers included complication rate, return of bowel function, number of consultations, reduction in total direct patient costs, and bed days saved.

Conclusions Our previous study showed that anesthesiologists can function as perioperative hospitalists who provide appropriate medical management while improving patient recovery. With this study, the impact of PHS remained up to one year after the transition of the co-management daily rounding service to a consultation only service. In fact, some patient categories showed further improvement in post-operative outcomes. These findings suggest that the transition of a daily rounding co-management PHS to a consultation only service is feasible without negatively impacting patient care.

28 IMPACT ASSESSMENT OF THE TRANSITION OF A CO-MANAGEMENT PERIOPERATIVE HOSPITALISTS MODEL TO A CONSULTATION MODEL TOWARDS PATIENT OUTCOME MARKERS
J Shen*, L Gad, D Ramsingh, D Narke, B Austin, G Steer. Loma Linda University, Loma Linda, CA

Purpose of study Perioperative care has wide variability in quality with conflicting models. In 2015, the Departments of Anesthesiology and Urology implemented a perioperative hospitalist service (PHS), consisting of anesthesiology-trained physicians, to co-manage patients during their perioperative period. From 2015 to 2016, the PHS team demonstrated improvements in patient recovery markers, decrease in hospital length of stay, and a reduction in total patient costs. In 2017, the PHS was phased out, but the patient care protocols remained active for the urology service. This study sought to evaluate if the improvement in patient care remained with the transition of PHS from a daily rounding co-management structure to a consultation service.

Methods used PHS was formed of selected anesthesiologists who received training on the core competencies for hospitalist medicine and were to medically manage patients undergoing major urologic procedures (prostatectomy, cystectomy, and nephrectomy). Impact was assessed by comparisons of patient outcome markers between the last year of the co-management structure service to the year of the consultation only service. The primary outcome marker was a reduction in length of stay. Secondary outcome markers included complication rate, return of bowel function, number of consultations, reduction in total direct patient costs, and bed days saved.

Conclusions Our previous study showed that such partnerships due to concern for decreasing incoming transfers, few studies report the true effect of such collaboration. We hypothesized that such partnership would increase community hospitalist pediatric care in the patient’s own community. Although some academic hospitals are hesitant to form these partnerships due to concern for decreasing incoming transfers, several studies report the true effect of such collaboration. We hypothesized that such partnership would increase community hospitalist pediatric care in the patient’s own community. Although some academic hospitals are hesitant to form these partnerships due to concern for decreasing incoming transfers, few studies report the true effect of such collaboration.

Methods used In July 2018, we established a partnership between an urban tertiary care academic medical center and a not-for-profit 194-bed community hospital, located 37 miles away. The partnership involved 1) staffing the

29 BENEFITS OF A COMMUNITY HOSPITAL – ACADEMIC UNIVERSITY PEDIATRIC PARTNERSHIP
M Hamline*, K Forman, K Tran-Viet, B McCollough, L Donati, H McKnight, S Lakshminrusimha, DA Lubarsky. University of California Davis Health, Sacramento, CA; Adventist Health Lodi Memorial, Lodi, CA

Purpose of study Academic children’s hospitals increasingly partner with community hospitals to provide inpatient neonatal and pediatric care in the patient’s own community. Although some academic hospitals are hesitant to form these partnerships due to concern for decreasing incoming transfers, few studies report the true effect of such collaboration. We hypothesized that such partnership would decrease community hospitalist pediatric volume and reduce transfers but increase case mix index (CMI) of transferred patients.

Methods used In July 2018, we established a partnership between an urban tertiary care academic medical center and a not-for-profit 194-bed community hospital, located 37 miles away. The partnership involved 1) staffing the
Summary of results Data regarding the community hospital inpatient pediatric population and pediatric transfers to the academic medical center are shown below. The academic center’s contribution margin increased by 123% from pre-to post-partnership. The partnership was cost-neutral at the departmental level.

Conclusions Our data suggest that the community-academic partnership successfully increased admissions and reduced LOS and inter-facility transfers while increasing acuity of transfers. These partnerships promote appropriate inpatient care and are cost-effective to both partners.

Purpose of study CKD affects approximately 30 million U.S. adults. Nutrition management can slow CKD progression and delay or prevent end-stage renal disease, but only 10% of patients with CKD receive any medical nutrition therapy (MNT) before initiating dialysis. This study aimed to identify factors that influence timely access to MNT for patients with NDD CKD from the perspective of patients and providers.

Methods used Three anonymous surveys were distributed to adults with NDD CKD, registered dietitian nutritionists (RDNs) and medical providers via REDCap. Surveys were distributed via the NKF 2019 Spring Clinical Meeting app, via Academy email listservs for RDNs and NKF email listservs for patients and medical providers. Descriptive analyses and Fisher’s exact tests were conducted with R (version 3.5.3).

Summary of results 348 patients, 66 RDNs and 30 medical providers responded. Patients were more optimistic than providers that they could make lifestyle changes to reduce CKD complications, with 44% of patients strongly agreeing with this statement vs. 29% of RDNs and 13% of medical providers. There was strong concordance regarding patient interest, with about two-thirds of medical providers, RDNs and providers responding. Patients were more optimistic than providers that patients would be interested in an RDN referral. However, there were feasibility concerns. Two-thirds of RDNs and medical providers felt that there are not enough specialists, banks, and the health department. Interventions included extensive outreach and frequent face-to-face communication with patients to assist with primary care and medication management. The CCCT reassessed patients, established care plans, and provided education for self-management of chronic conditions.
enough RDNs with expertise in renal nutrition available for referrals. Around 40–50% of RDNs and medical providers felt that the EMR was not set up to make RDNs referrals easy, and that there was not adequate insurance coverage. About half of RDNs were aware of Medicare coverage for MNT, but few medical providers or patients were familiar with the coverage.

Conclusions Many patients with NDD CKD are interested in MNT and confident that it can help them with disease management. Perceived referral and reimbursement barriers should be further explored.

Purpose of study Effective analgesic modalities improve outcomes in patients undergoing total knee arthroplasty (TKA). We hypothesized implementation of a relatively new nerve block method, the iPACK block would be associated with decreased length of stay and 24-hour opioid consumption in TKA.

Methods used This retrospective cohort studies outcomes of interest were postoperative length of stay (LOS) and postoperative 24-hour oral morphine equivalent (OME) consumption. All patients received a single shot adductor canal block (ACB) and local infiltration analgesia (LIA) and were grouped based on whether they received an iPACK block. Potentially confounding variables such as age, gender, BMI, presence of comorbid conditions, operation room time, and current chronic steroid or opioid medication therapy were recorded. Descriptive statistics characterized patients in our sample. Categorical demographic and clinical group differences were examined using Chi-square analysis. Continuous group differences were explored using 2 sample t-tests for symmetrical distributions and Mann-Whitney U for skewed distributions. Linear regression models were performed to detect differences in study outcomes between the two groups, controlling for confounding variables.

Summary of results Incorporation of the iPACK block was not associated with reduced LOS unadjusted (p=0.11) or in fully controlled models (p=0.21). Potentially modifiable factors that were found to increase length of stay included chronic opioid consumption (p=0.04) and case duration (p=0.01). Use of the iPACK block was not associated with a change in 24-hour OME consumption unadjusted (p=0.52) or in fully controlled models (p=0.34). Potentially modifiable factors that significantly increased opioid consumption were a history of chronic opioid therapy (p=0.002) or corticosteroid therapy (p=0.003).

Conclusions As only the second study performed investigating incorporation of an iPACK into a pathway already including LIA and ACB we have contributed to a small body of evidence suggesting the iPACK block may not add significant benefit alongside these methods. To date no prospective study has investigated outcomes associated with lone addition of iPACK block into a pathway including ACB and LIA. Further randomized research is warranted.

Conclusions Point-of-care ultrasound (POCUS) has rapidly emerged as a modality to improve bedside assessment. The driving force for this adoption is that clinical decision-making
can be improved over traditional physical examination with the application of POCUS. Since the limiting factor for the adoption of POCUS is education, the authors launched a free open access educational platform. We present our initial data from our online users on improving content knowledge.

**Methods used** Over a two-year period a validated educational POCUS curriculum, the FORESIGHT Examination, was transitioned to an online platform (www.foresightultrasound.com). The curriculum was developed to educate and evaluate: cardiovascular function, volume status, pulmonary function, presence of abdominal fluid, gastric content, position of the endotracheal tube, and presence of increased intracranial pressure. Performance of the website was evaluated via surveys on the content and content multiple choice testing. The primary outcome marker was improvement in content testing scores after reviewing the online curriculum. Analysis was performed via student t-test, a p value of significance equal to 0.05.

**Summary of results** To date 530 users have accessed the website. Survey results on the website content to improve POCUS knowledge showed that 83% reported to strongly agree, 10% to agree, and 7% were neutral. Review of content testing showed a statistically significant (p<0.05) improvement in scores from pre to post training for each testing category: 1. ultrasound physics, 2. mechanisms of hypotension/volume status, 3. cardiac, 4. pulmonary, and 5. abdominal/vascular access/neuro.

**Conclusions** While there is still much to be developed, the concept of P-POCUS (perioperative POCUS) has shown dramatic growth in recent years. Further innovation and integration of this tool have potential for providing the quality of care we can provide at the bedside. This study highlights one successful platform to improve knowledge on this topic.

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**WHAT MATTERS MOST: AN INTERVENTION FOR LEARNERS**

SD Andersen*, H Lokos, A Hayton. Loma Linda University, Loma Linda, CA

10.1136/jim-2019-WMRC.35

**Purpose of study** Inquiring about what matters most has been deemed a powerful tool to support greater patient wellbeing. The purpose of this study was to better equip junior medical students to offer whole person care (WPC). We expect that perceived value, confidence and frequency of students asking patients what matters most to them will increase after the intervention. We hope as a result of students being more engaged with their patients, they will feel more involved on the team.

**Methods used** A pre-baseline survey was administered to junior medical students at the beginning of their Internal Medicine clerkship. A workshop was conducted early in the clerkship. Students were given a pocket guide with questions aiding connection with patients. Students were told to ask each patient they saw about what matters to them and present this information on rounds. Attendings and residents were encouraged by email to encourage students to present. A post-baseline survey was issued to each group at the end of the clerkship. The main barriers selected by students were time constraints, workload, and lack of support. Students reported that the intervention increased sense of meaning. Students perceived a need for more role models and earlier training in their pre-clinical years to enhance the effectiveness of this intervention.

**Conclusions** Preliminary results show that students express a significant increase in the value they place on WPC after undergoing the intervention. We anticipate that as the study’s power increases with more student responses, we will see a statistically significant difference in student confidence in ability to provide WPC. As students engage on this level with their patients we expect they will find more meaning in their practice.

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**Immunology and rheumatology I**

**Concurrent session**

**12:45 PM**

**Thursday, January 23, 2020**

**36**  **SECUKINUMAB AND INCIDENT INFLAMMATORY BOWEL DISEASE IN ANKYLOSING SPONDYLITIS**

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10.1136/jim-2019-WMRC.36

**Purpose of study** Secukinumab is a fully human monoclonal antibody against interleukin-17A for use in treating active ankylosing spondylitis (AS). Inflammatory bowel disease (IBD) is increased in AS and subclinical gut inflammation is found in 50–60 percent of AS patients. Incident IBD and exacerbation of established IBD was reported as an adverse event in secukinumab clinical trials, though these were rare events (<1%) and not significantly different compared to the placebo population. There have been 60 reported cases of IBD, including ulcerative colitis, and Crohn’s disease in the FDA Adverse Events Reporting System (FAERS) in AS patients treated with secukinumab, with 2 cases listed as exacerbation of preexisting disease. We evaluated the incidence of IBD in AS patients after initiation of secukinumab.

**Methods used** Data was extracted from a prospective axial spondyloarthritis (axSpA) cohort followed by a tertiary care center in a subspecialty spondyloarthitis clinic at the University of California, San Francisco. We describe a case-series of three AS patients who developed incident IBD after treatment with secukinumab. The patient demographics, treatment exposures, medication use and disease course since initial presentation at the clinic are reported.

**Summary of results** In our axSpA cohort, 22 patients with AS were treated with secukinumab. Three patients (13.6%) developed incident IBD (2 Crohn’s disease and 1 ulcerative colitis). The patients experienced symptoms of IBD within seven months (1.5–7) after starting secukinumab and were...
referred to a gastroenterologist in the setting of new diarrhea. IBD diagnoses were confirmed by a gastroenterologist in the setting of a clinical evaluation, colonoscopy and histopathology.

**Conclusions** Our findings suggest that IBD incidence may occur more frequently with secukinumab exposure than has been reported. There are limitations to this study as it is uncontrolled, and the patient population may have more severe disease at a tertiary care center. We recommend that providers treating AS patients with secukinumab take a detailed gastrointestinal history and have a low threshold for further evaluation in those with new gastrointestinal symptoms.

**37 SECONDARY EFFICACY FAILURE IS THE PREDOMINANT ABSTRACT WITHDRAWN HAIR REPIGMENTATION AFTER RITUXIMAB USE IN A CORRELATION BETWEEN CLINICAL AND IN VITRO GENETICITY OF TNFi, IN EFFORT TO INCREASE PERSISTENCE AND IMMUNOCLOGICAL DISORDERS**

**Purpose of study** Psoriatic arthritis is a chronic inflammatory disease of the joints and skin that affects 1/1000 people in the US. Tumor necrosis factor inhibitors (TNFi) are used when symptoms are severe. However, current research describing TNFi persistence rates, defined as time from initiation to discontinuation of the drug, is inconsistent and incomplete. This study examined characteristics associated with persistence of TNFi (adalimumab, certolizumab pegol, etanercept, golimumab, and infliximab) and reasons for discontinuation.

**Methods used** US veterans enrolled in the Program to Understand the Longterm Outcomes in Spondyloarthritis (PULSAR) from 2007 – 2017 who 1) were diagnosed with psoriatic arthritis or psoriasis and 2) had been treated with a TNFi were included in the study. Stata was used to conduct Student’s t-tests, Pearson Chi squared tests, time-to-event analyses, and multivariate analyses.

**Summary of results** 321 individuals with 931 TNFi courses were included in the study. The mean age was 55.4 years, and 83.8% of the cohort continued at least one TNFi course at one year. Course order was correlated with persistence on a TNFi at one year (HR 1.09, p<0.01). Baseline Bath Ankylosing Spondylitis Functional Index (BASFI) and baseline pain scores were correlated with response to a TNFi (HR 1.12, p=0.018 and HR 0.689, p<0.01, respectively). Other demographics were not independent predictors of response or persistence. Infliximab (79.3%) was associated with improved persistence at 24 months when compared to adalimumab (66.4%, p=0.019) and certolizumab (54.1%, p=0.013). The most commonly cited reason for discontinuing TNFi treatment was secondary failure (33%), defined as a loss of prior efficacy after >6 months of treatment, followed by adverse events (25%).

**Conclusions** This study found that most clinical characteristics, other than course order, did not affect persistence on a TNFi, suggesting that persistence is difficult to anticipate. Since persistence is suboptimal, further research should study possible mechanisms to explain loss of efficacy, including the immunogenicity of TNFi, in an effort to increase persistence and improve treatment outcomes.

**38 CORRELATION BETWEEN CLINICAL AND IN VITRO RESPONSE TO METHOTREXATE IN RHEUMATOID ARTHRITIS**

**Purpose of study** Rheumatoid arthritis (RA) is a systemic autoimmune disease that decreases quality of life and can lead to disability. Methotrexate (MTX), which targets activated T cells, is the first-line of therapy for RA. Unfortunately, 30 to 50% of RA patients do not experience adequate reduction of symptoms while on MTX. Predicting MTX response would be helpful for guiding treatment decisions and improving the lives of RA patients. The purpose of this pilot study was to explore a possible association between clinical response to MTX and in vitro effects of MTX on leukocyte secretion of IL-17 and IFN-g, pro-inflammatory cytokines associated with the activated Th17 and Th1 cells.

**Methods used** Peripheral blood samples were collected from 16 RA patients prescribed MTX but prior to initiation of MTX treatment. Leukocytes were isolated from the blood samples, activated with anti-CD3 and anti-CD28 antibodies, and treated with several concentrations of MTX (0 to 1 uM). Supernatant was harvested at 72 hours, and concentrations of IL-17 and IFN-g were assessed by ELISA. The RA patients’ charts were reviewed to assess the clinical effects of MTX after 3 months of MTX treatment, as described by changes in clinical disease activity index (CDAI) and any side effects.

**Summary of results** In most blood samples, 1 uM of MTX decreased secretion of IL-17 and IFN-g. Maximal decrease was -28 - 87% for IL-17, and 15 – 92% for IFN-g. The concentration of MTX required to achieve half-maximal decrease (IC50) also varied for both IL-17 (0.02 – 0.3 uM) and IFN-g (0.01 – 0.3 uM). Based on samples from 9 patients, there is greater variation in the IL-17 IC50s of those with side effects vs those without (p<0.05), with absence of side effects associated with IL17 1C50 0.03 – 0.05 uM.

**Conclusions** There is large person-to-person variation in the in vitro effects of MTX on IL-17. Increased sample size is needed to assess whether this variation could be of clinical significance.

**39 ABSTRACT WITHDRAWN**

**40 HAIR REPIGMENTATION AFTER RITUXIMAB USE IN A PATIENT WITH SYSTEMIC SCLEROSIS**

**Purpose of study** Repigmentation of gray/white hair is rarely reported in the literature. There are several etiologies contributing to this, and autoimmune diseases have been considered one of them. An overactive immune response could result in sudden graying of hair in mouse models through inducible expression of melanogenesis associated transcription factor (MITF), which controls the production and life cycle of melanocytes and acts as an off-switch to innate immune target.
genes at the same time. Similarly, change of hair color have been associated with a multitude of drugs including corticosteroids, cyclosporine and most recently targeted anticancer therapies targeting immune system. However, very little is known of the effects of biologics used in autoimmune diseases and their role in hair repigmentation.

Methods used A 70 year old male with systemic sclerosis was started on Rituximab therapy in 09/2017, now s/p 5 cycles of Rituximab given every 6 months. Initially presented with Raynaud’s, shortness of breath, weight loss, arthralgias and skin findings. On presentation, physical examination revealed achromotrichia (silver hair) of normal aging, without hair loss or thinning. After 6–9 months into Rituximab therapy, patient had repigmentation of hair to black. Now he has over 60% of repigmentation and is in third year of rituximab therapy with stable disease.

Summary of results Graying of hair usually begins in the third and fourth decade of life after melanocyte depletion unless there are other factors including nutritional deficiencies, stress, and smoking among others. In those instances, hair changes can be seen earlier. Rituximab among other biologics has been used in several autoimmune conditions including SLE, rheumatoid arthritis and systemic sclerosis. Alopecia is considered a side effect commonly seen, but hair repigmentation is not reported. To our knowledge this is the first ever reported case of repigmentation in a patient on Rituximab for systemic sclerosis. Hair darkening after use of biologics might be reversible but the exact mechanism remains elusive.

Conclusions To our knowledge this is the first ever reported case of repigmentation in a patient on Rituximab for systemic sclerosis. Hair darkening after use of biologics might be reversible but the exact mechanism remains elusive.
Conclusions Medical bias can increase the difficulty in distinguishing between sepsis and an acute episode of SLE can be challenging in a patient without prior history of medical illness. Therefore, clinicians should have a high degree of suspicion for autoimmune diseases (i.e., SLE) when a patient presents with a life-threatening and multiorgan condition.

Infectious diseases I
Concurrent session
12:45 PM
Thursday, January 23, 2020

**EFFECT OF EXTRACORPOREAL SHOCKWAVE ON RHIZOPUS ORYZAE BIOFILM**

1KH Anderson*, 2P Sksek, 3K Olga, 4T Hillick. 1University of Utah, Salt Lake City, UT; 2Ludwig Boltzmann Institute for Experimental and Clinical Traumatology, Vienna, Austria; 3Utah Valley University, Orem, UT

**Purpose of study** Fungal biofilms are communities of adherent cells surrounded by an extracellular matrix. Clinically, biofilm associated infections can be extremely difficult to eradicate due to their resistance to antifungals and host defense mechanisms. In this study we investigate the effect of extracorporeal shockwave treatment (ESWT) on Rhizopus oryzae biofilm.

**Methods used** Biofilms were treated under four main conditions: Shock only, Shock + Amp B, Amp B control and Biofilm control. Shocks were applied to pre-biofilm, 12, and 24 hours post plating (mature biofilm). Tubes were exposed to 300 pulses using an energy density of 0.55 mJ/mm² at 3 Hz. Shock wave treatment was performed using a water bath setup.

**Summary of results** Results indicate that shockwave application in combination with antifungal treatment increases fungal cellular respiration, extracellular matrix production and hyphal width compared to antifungal treatment alone. Shockwave when applied in isolation, (without antifungal treatment) revealed no difference in respiration or matrix production.

**Conclusions** It is suggested that shockwaves disrupt biofilm structure allowing antifungal drugs to penetrate and interact with the underlying cells causing an increase in cellular respiration, hyphal growth and extracellular matrix production. This could allow the host system to recognize pathogen proteins and increase pathogen clearing by the immune system. However, further research will need to be done to confirm this hypothesis.

**CONGENITAL SYphilIS: A QUALITATIVE RETROSPECTIVE LONGITUDINAL STUDY ON CHARACTERISTICS OF MOTHERS AND NEWBORNS IN A TERTIARY CARE HOSPITAL**

R Kaushal*, H Chow, R Jain, C Nelson, C Rongkavilit. UCSF Fresno, Fresno, CA

**Purpose of study**
1. To determine maternal factors leading to increased prevalence of congenital syphilis in our community and state.
2. To study the effects in newborns of in utero exposure to untreated or partially treated maternal syphilis.

**Methods used** ICD-9 and 10 codes were used to identify newborns born to mothers with a diagnosis of syphilis at the time of delivery at a tertiary care hospital in Fresno, California from 01/2010 -12/2018. 231 newborns and mother dyads were included in analysis.

**Summary of results**
Maternal characteristics: The mean age was 27 years old (range 16-42 years). 57 (24.7%) were Caucasian and 167 (75.3%) were ethnic minorities. 203 (88.7%) were unmarried, 207 (89.6%) were unemployed. 17 (7.4%) were incarcerated, 18 (7.8%) were victims of domestic violence. 136 (58.9%) were admitted to drug use, of which 112 (48.5%) used methamphetamines, 57 (24.7%) used marijuana, and 15 (6.5%) used opiates. 78 (33.8%) smoked tobacco, 14 (6.1%) drank alcohol during pregnancy. 61 (26.4%) positive for HSV by serology. 35 (15.2%) positive for Chlamydia, 11 (4.8%) positive for Gonorrhea, 6 (2.6%) positive for HIV. 75 (32.5%) were treated adequately for syphilis during pregnancy, 119 (51.5%) treated inadequately, and 34 (14.7%) received no treatment.

Newborn characteristics: Of 231, 67 were born from 2010–14 (20.4%) and 184 from 2015–18 (79.6%). At delivery, 19 (8.2%) newborns were a gestational age≤32 weeks, 57 (24.7%) 33–37 weeks, and 155 (67.1%) >37 weeks. 15 (6.5%) had a positive CSF VDRL, 36 (15.6%) had an abnormal bone survey, 5 (2.2%) had an abnormal eye examination. 220 (95.2%) received treatment for syphilis, and 7 (3.0%) did not receive treatment. Of those treated, 56 (24.2%) received IM Penicillin, 139 (60.2%) received IV Penicillin, and 28 (12.1%) received both IM and IV Penicillin.

**Conclusions** We identified that ethnic minorities, unemployment, illicit drug use, STDs were prevalent among pregnant women with syphilis. High rates of inadequate/no treatment during pregnancy is concerning and have led to prolonged hospitalization of newborns for IV therapy. Strengthening public health infrastructures to test and care for these high-risk pregnant women are urgently needed.

**COMPARISON OF PROCALCITONIN AND C-REACTIVE PROTEIN (CRP) IN NEONATAL BACTERIAL SEPSIS**

1A Bilg*, M Choi, S Tewari, B Chan, S Anis, J Luu, B Afghani. 1UC Irvine School of Medicine, Irvine, CA; 2UC Riverside School of Medicine, Riverside, CA; 3CHOC Hospital of Orange County, Orange, CA

**Purpose of study** Blood culture, the gold standard for the confirmation of bacterial sepsis, has significant limitations including time delay in obtaining results. Biomarkers such as serum
Abstracts

Abstract 45 Table 1 Comparison of PCT & CRP in early-onset neonatal sepsis

<table>
<thead>
<tr>
<th>First Author, Year</th>
<th>Study Subgroups</th>
<th>Test Threshold</th>
<th>Test Sensitivity</th>
<th>Test Specificity</th>
</tr>
</thead>
<tbody>
<tr>
<td>Guibourdenche 2002</td>
<td>21/120 confirmed sepsis or probable infection, 88/120 bacterial culture-negative, 11/120 discordant clinical and bacteriological findings</td>
<td>PCT 2.5 μg/L</td>
<td>PCT 87%</td>
<td>PCT 90%</td>
</tr>
<tr>
<td>Groselj-Grenc 2009</td>
<td>17/46 SIRS with bacterial sepsis, 29/46 SIRS without bacterial sepsis</td>
<td>CRP 7.5 mg/L</td>
<td>CRP 68%</td>
<td>CRP 80%</td>
</tr>
<tr>
<td>Altunhan 2011</td>
<td>171/260 positive culture and clinical signs of sepsis, 89/260 no clinical signs of sepsis or risk factors</td>
<td>PCT 2.28 μg/L</td>
<td>PCT 82%</td>
<td>PCT 48%</td>
</tr>
<tr>
<td>Mohsen 2015</td>
<td>35/70 positive blood culture and clinical sepsis, 35/70 healthy controls</td>
<td>PCT 11 mg/L</td>
<td>PCT 59%</td>
<td>PCT 100%</td>
</tr>
<tr>
<td>Al-Zahrani 2015</td>
<td>30/100 positive blood culture and/or positive bacterial 16S rDNA PCR, 37/100 negative blood culture and positive sepsis clinical symptoms and screen, 29/100 negative blood culture and screen</td>
<td>CRP 5.9 mg/L</td>
<td>CRP 45.5%</td>
<td>CRP 59.4%</td>
</tr>
<tr>
<td>Elmouttaleb 2016</td>
<td>50/80 clinical sepsis and positive blood culture, 30/80 healthy controls</td>
<td>PCT 1.1 pg/mL</td>
<td>PCT 80.0%</td>
<td>PCT 85.7%</td>
</tr>
</tbody>
</table>

Early onset=within 3 days of life, SIRS=systemic inflammatory response syndrome

Abstract 46 Figure 1

CHARACTERIZATION OF HIV QUASISPECIES VIA PARTICLE-TEMPLATED EMULSIFICATION

WY Shin*, M Hatori. UCSF, San Francisco, CA

Purpose of study RNA retroviruses can form ‘similar-but-different’ variants (quasispecies). These variants are found through next-generation sequencing to track shifts in quasispecies during early infection. A disadvantage is that it may be expensive to sequence a long stretch of RNA if needed. Droplet microfluidic devices create micron size water-in-oil droplets, where rare cells or novel genes can be trapped into one droplet and analyzed. The advantages are that it is high-throughput and uses less reagents. Barriers to microfluidics are the hardware, expertise, and cost needed to fabricate the devices. The goal of this project is to see whether the method particle templated emulsification (PTE) can be used to track rare HIV RNA variants. This method creates droplets simply by vortexing polyacrylamide beads with your sample. We also want to show that we can do Splicing by Overlap Extension (SOE) PCR in PTE, which fuses two separate pieces of DNA to reduce the size of the amplicon.

Methods used Droplet microfluidic devices are made via standard photolithography. Polyacrylamide beads for PTE are made via an air-triggering device that splits the flow of acrylamide into micron size beads. We mix the PCR reagents with the polyacrylamide beads and allow them to incubate. For PTE, we vortex the mixed beads in a solution of surfactant. This process creates an emulsion encapsulating each bead with the PCR reagents. The beads are transferred to PCR tubes and run through a standard PCR program. The amplified products in the beads are viewed under a microscope.

Summary of results After PCR amplification, we see that the control group and SOE PCR reaction have amplified product. However, we get the incorrect size product for the SOE PCR reaction, meaning SOE PCR did not work in PTE.

Conclusions Further optimization is needed for SOE PCR to work in PTE. We will use a different test plasmid that encodes the HIV gag gene. We would repeat the PTE experiments again with the new plasmid and further optimize the protocols. If there is no progress, one alternative to the project is to use primer studded beads as the template for PTE with unique molecular identifiers.
INTERFERON BETA INCREASED SIGLEC-1 EXPRESSION ON HUMAN GUT MACROPHAGES

SM Lada*, CM Purba, SM Dillon, ML Santiago, CC Wilson. University of Colorado Anschutz Medical Campus, Aurora, CO
10.1136/jim-2019-WMRC.47

Purpose of study A breakdown in gut homeostasis is a key feature of HIV-1 infection, in part due to the high levels of HIV-1 replication that occur in early in the gut. Type 1 Interferons (IFN-1s) are innate cytokines that play critical roles in controlling viral replication. Siglec-1 (CD169) expression, induced by IFN-1s on myeloid dendritic cells (mDC) and macrophages (MΦ), binds HIV and has been implicated in HIV trans-infection of CD4 T cells. Increased CD169 in response to gram-negative (GN) bacterial cell wall component lipopolysaccharide (LPS) has also been reported. We previously showed elevated levels of the IFN-1 interferon beta (IFNβ) as well as LPS in gut tissue of people living HIV. CD169 is expressed on murine gut MΦ, but few studies have investigated expression in human gut tissues or its regulation. We hypothesized that exposure to IFNβ or GN bacterial products would increase CD169 expression on human gut MΦ.

Methods used Human jejunal lamina propria mononuclear cells (LPMC), obtained from healthy discarded surgical tissue (N=4), were cultured for 18 hrs with IFNβ (1000–1 pg/mL; 10-fold dilutions), enteric commensal GN Escherichia coli lysate (10ug/mL), or were unstimulated. Flow cytometry was used to evaluate expression of CD169 on MΦ or mDC pre (baseline) and post in vitro culture.

Summary of results MΦ were identified as HLA-DR+CD64+CD11c± cells within viable CD45+CD3-CD19- LPMC. At baseline MΦ constituted 0.16±0.03% (Mean±SEM); of these MΦ 7.95±4.1% expressed CD169. HLA-DR+CD64+CD11c+ mDC within viable CD45+CD3-CD19- LPMC, constituted 0.11±0.02% of baseline; there were few mDC expressing CD169. In the presence of 1000 pg/mL IFNβ, 27±4% of MΦ expressed CD169, a 6.2-fold increase over unstimulated. With 100 pg/mL IFNβ, 20±5% of MΦ were CD169+, a 4.2-fold increase over unstimulated. At lower doses of IFNβ, CD169 expression by MΦ only increased by 1.5-fold versus unstimulated. Exposure to E. coli lysates increased CD169 expression on MΦ by 2.3-fold.

Conclusions IFNβ induced CD169 expression on human gut MΦ in vitro in a dose dependent manner. Additionally, GN commensal bacteria lysates increased expression, although to a lesser degree than high doses of IFNβ. Future studies will investigate the role of CD169+ gut MΦ in T cell HIV infection.

IMPLEMENTING A PEDIATRIC ANTIBIOTIC STEWARDSHIP PROGRAM IN A MULTIDISCIPLINARY PRIMARY CARE CLINIC

CH Kim*, S Goggin, y Garcia E Fernandez, N Nakra, U Shaikh. UC Davis Health, Sacramento, CA
10.1136/jim-2019-WMRC.48

Purpose of study Family practitioners, nurse practitioners, and physician assistants prescribe higher rates of antibiotics for acute respiratory infections (Agrio 2018). We established a pediatric resident teaching clinic within a local primary care center where pediatric care had previously been delivered by clinicians without pediatric training. Guided by the CDC’s ‘Core elements of outpatient antibiotic stewardship,’ we identified guideline-based pediatric antibiotic prescribing as an area for quality improvement (QI). We aimed to reduce unnecessary antibiotic prescriptions (Rx’s) for viral diagnoses by 15% and inappropriate Rx’s (based on drug choice, dose, or duration) by 25% for patients ≤11 years old over an 18-month intervention period.

Methods used We implemented regular feedback sessions on Rx rates, as well as treatment guideline review sessions. We used 3-month PDSA cycles to identify ongoing opportunities for improvement. Pediatric residents performed chart reviews for systemic antibiotic Rx’s identified in the EMR. We determined the proportion of antibiotic Rx’s that were unnecessary or inappropriate. We compared data from the intervention period to a 10-month baseline period.

Summary of results During the first 8 months thus far, compared to baseline, the proportion of unnecessary Rx’s decreased by 10% (27% vs 17%) and inappropriate Rx’s by 47% (82% vs 35%). Interventions performed included a baseline session at month 0, review of acute otitis media guidelines at month 1, and feedback session and review of streptococcal pharyngitis guidelines at month 4. From month 5 through month 8, an EMR change led to a delay in obtaining charts for abstraction, inability to provide feedback, and a rise in inappropriate Rx’s (24% vs 35% mo. 8), though unnecessary Rx’s continued to decline during this period (21% mo. 4 vs 17% mo. 8).

Conclusions Utilizing QI techniques in the outpatient setting to improve adherence to pediatric antibiotic prescribing guidelines has yielded promising early results, though technical challenges have reinforced that ongoing interventions are needed to sustain these results. We identified streamlining of data collection and providing individualized, real-time feedback with behavioral nudges (Meeker 2016, Linder 2017) to clinicians as areas for future improvement.

MACROPHAGE SUPPRESSION DURING THE ACUTE PHASE OF CONGENITAL CYTOMEGALOVIRUS INFECTION DOES NOT CHANGE HEARING OUTCOMES IN A MOUSE MODEL

10.1136/jim-2019-WMRC.49

Purpose of study To determine whether treatment with clodronate-containing liposomes reduces sensorineural hearing loss (SNHL) in cytomegalovirus-infected mice and to evaluate the effect of clodronate-containing liposomes on systemic macrophage count.

Methods used BALB/c mice were inoculated intracranially with murine-CMV (mCMV) or saline on postnatal day 3. The mice were treated with intraperitoneal injections of clodronate-containing or control liposomes over 14 days. Auditory thresholds were assessed using Distortion Product Otoacoustic Emission (DPOAE) and Auditory Brainstem Response (ABR) testing at three weeks of age. Spleens were harvested at 9 days post-infection and evaluated using flow cytometry to confirm global knockdown of monocyte macrophages by clodronate-containing liposomes.
Summary of results MCMV-infected mice treated with clodronate-containing liposomes did not show significant reduction in ABR ($P>0.5$ by Kruskal-Wallis test) and DPOAE ($P>0.5$) thresholds compared to mCMV-infected mice treated with control liposomes. Significant differences were observed in ABR and DPOAE thresholds of mCMV-infected mice compared to non-infected mice ($P<0.001$). MCMV-infected mice treated with clodronate-containing liposomes showed significant reduction of splenic monocyte macrophages compared to mCMV-infected mice treated with control liposomes ($P<0.01$ by t-test).

Conclusions Treatment with clodronate-containing liposomes did not change hearing thresholds in mCMV-infected mice at three weeks of age although clodronate treatment did cause a global knockdown of monocyte macrophages. These results warrant further investigation into the role of specific subpopulations of cochlear macrophages in the development of progressive SNHL in congenital CMV infection.

**TYROSINE KINASE INHIBITION TO REDUCE THE HIV-1 RESERVOIR**

M Szaniawski*, A Spivak. University of Utah, Salt Lake City, UT

10.1136/jim-2019-WMRC.50

**Purpose of study** To validate the efficacy of tyrosine kinase inhibitors (TKI) as anti-proliferative and anti-HIV-1 agents in CD4+ T cells and myeloid cells, respectively. Ultimately, these data will inform the design of a clinical trial focused on the use of TKI as an HIV-1 cure strategy.

**Methods used** Whole blood was obtained by peripheral phlebotomy from healthy donors, and peripheral blood mononuclear cells (PBMC) were isolated followed by purification of CD14+ monocytes and CD4+ memory T cells. Monocytes were differentiated over 7 days in autologous serum to generate monocyte-derived macrophages (MDM).

Summary of results TKIs inhibit proliferation induced by IL-2, IL-7, or IL-15 in memory CD4+ T cells in vitro, preventing the major mechanism responsible for expansion and persistence of the viral reservoir. We find that HIV-1 infection induces a senescent phenotype in macrophages, which is further counteracted by the use of dasatinib. Finally, TKIs are potent inhibitors of HIV-1 in myeloid cells through activation of the host restriction factor SAMHD1, which prevents new infection through direct inhibition of reverse transcription.

**Conclusions** We conclude that dasatinib possess two major immunomodulatory qualities that make them attractive candidates for HIV-1 cure. As an anti-proliferative, dasatinib interrupts homeostatic proliferation, the major mechanism underlying HIV-1 persistence in CD4+ T cells. In macrophages, dasatinib both directly inhibits HIV-1 reverse transcription and counteracts HIV-1 infection-induced cellular senescence.
transcription through the activation of SAMHD1 and counteracts the pro-inflammatory effect of HIV-1 infection in myeloid cells, a process that may underlie HIV-1 comorbidities including cardiovascular disease (CVD) and HIV-associated neurocognitive disease (HAND).

**Abstracts**

**51 PATHOGEN AND ANTIMICROBIAL RESISTANCE SURVEILLANCE IN UGANDAN HIV POSITIVE ADULTS WITH PNEUMONIA**

1JD Bloomstein*, 2S Caldera, 2Z Huang, 3W Worodria, 3B Byanyima, 2E Musisi, 5S Kaswabuli, 3J Zawedde, 2J Moore, 2S Lynch, 3M Shenyu, 2C Langelier, University of California, Davis, Sacramento, CA; 2University of California, San Francisco, San Francisco, CA; 3Makarere University, Kampala, Uganda

**Purpose of study** Preliminary analyses using machine learning algorithms developed to predict phenotypic antimicrobial resistance from high dimensional genomic data demonstrate 85–100% concordance between culture-determined resistance and detection of antimicrobial-specific resistance genes from cultured isolates. To further evaluate the algorithms, Next Generation Sequencing (NGS) data from Ugandan HIV positive adults with pneumonia was analyzed to identify the type of infection- bacterial pneumonia, viral pneumonia, presence of TB, and disease correlations with CD4 count.

**Methods used** We implemented the Rules-Based Method to identify individual or multiple pathogens implicated in each pneumonia case for 218 patients. Using RNA-seq analysis of patient samples, we first analyzed the landscape of the patient virome and determined which patients had active CMV or EBV infections. Next we evaluated for the presence of Pneumocystis jirovecii infection. Subsequently we identified patients with Mycobacterium tuberculosis infections by finding an optimal threshold of sequencing reads that maximized sensitivity.

**Summary of results** Preliminary bacterial results show that the most common pathogen were Haemophilus influenzae, Pseudomonas aeruginosa, and Pneumocystis jirovecii. Preliminary viral results show that the most common secondary viruses were Rhinovirus and Influenza. While all 8 patients positive for Pneumocystis jirovecii bronchoalveolar Giemsa stain were positive by NGS, 2 patients had high read numbers by NGS.

**Conclusions** The RBM showed high sensitivity for detection of bacterial, viral, and fungal pathogens in pneumonia patients. NGS showed high sensitivity for Pneumocystis jirovecii in particular. Preliminary correlations between infection with CMV, and other viruses, with CD4 count <200 suggest that there could be a potential connection between immunosuppression due to AIDS and certain viruses that has not been previously appreciated. The potentially high prevalence of Pseudomonas aeruginosa could have important consequences for clinical care. Current empirical treatment at the hospital is Ceftriaxone, which does not cover Pseudomonas aeruginosa.

**Neonatology general I**

**Concurrent session**

**12:45 PM**

**Thursday, January 23, 2020**

**52 EFFECT OF CONTACT PRECAUTIONS ON STAPHYLOCOCCUS AUREUS AND CLINICAL OUTCOMES OF COLONIZED PATIENTS IN THE NEONATAL INTENSIVE CARE UNIT**

5S Shahbaz*, M Garg, A de St Maurice, M Sim. UCLA, Los Angeles, CA

**Purpose of study** The primary objective is to assess the incidence of S. aureus colonization and invasive disease during and after contact precautions. The secondary objective is to evaluate clinical outcomes of colonized patients versus non-colonized controls.

**Methods used** A retrospective chart review of all infants at two UCLA NICUs who screened positive for S. aureus (n=83). Information was also collected for controls with negative screens (n=151). Of note, from Aug 2014 to Aug 2016 contact precautions were mandated; they were universally discontinued from Sep 2016 to November 2018.

**Summary of results** There was a lower incidence of S. aureus colonization when contact precautions were utilized (28.6% vs 44.6%, p=0.01). Of the positively colonized patients, 20.5% were colonized with MRSA and 79.5% with MSSA. There were twice as many positive invasive MSSA cultures than MRSA. Most invasive cultures occurred in previously colonized infants (100% for MRSA, 73.9% for MSSA). There was a higher rate of invasion with MRSA (41.2%) than MSSA (16.7%). There were no differences in measured clinical outcomes in colonized patients before and after contact precautions (e.g., duration invasive mechanical ventilation, incidence of chronic lung disease and necrotizing enterocolitis, NICU length of stay and in-hospital mortality). Colonized patients were 38.5 more likely to be associated with a soft tissue infection (p=0.0005) and have an average 27 day longer NICU length of stay (p<0.001) when compared to noncolonized patients (when controlling for gestational age and surgical status).

**Conclusions** There was a lower incidence of S. aureus colonization when contact precautions were utilized. Most invasive cultures occurred in colonized patients, with a higher rate of invasion for MRSA. There was a higher incidence of MSSA colonization and invasive cultures. There were no differences in clinical outcomes among colonized patients during and after contact precautions. Colonization was more likely to be associated with soft tissue infection and longer NICU length of stay.
ASSOCIATIONS BETWEEN MATERNAL PRE-PREGNANCY OBESITY AND NEONATAL NEUROBEHAVIOR IN INFANTS BORN BEFORE 30 WEEKS GESTATION

N Nosavan*, L Smith, J Hofheimer, E McGowan, T O'Shea, S Pastynak, C Neal, B Carter, J Helderman, J Check, A Sollman, M Roberts, L Dansereau, S DellaGrotta, B Lester. The NOVI Research Network, Providence, RI

10.1136/jim-2019-WMRC.53

Purpose of study The rate of pre-pregnancy obesity has steadily increased, increasing the risk for preterm delivery, gestational diabetes, pregnancy-induced hypertension, and cesarean sections. Elevated pre-pregnancy weight has also been linked to poor long-term child neurodevelopmental outcomes such as decreased cognitive performance, behavioral and emotional difficulties, and ADHD symptoms. The purpose of our study is to examine the relationship between maternal pre-pregnancy obesity and short-term neonatal neurobehavior in infants born <30 weeks post-menstrual age.

Methods used The Neonatal Neurobehavior and Outcomes in Very Preterm Infants (NOVI) study recruited at 9 NICUs affiliated with 6 universities participating in the Vermont-Oxford Network. The NICU Network Neurobehavioral Scale (NNNS) was used to assess neurobehavior at discharge.

Summary of results 709 infants enrolled in NOVI and 664 had completed neurobehavioral assessments and medical data. Of those 227 (34.2%) infants were born to mothers with pre-pregnancy obesity (BMI >30). In unadjusted analyses, stress abstinence and non-optimal reflexes on the NNNS were higher in infants born to mother with pre-pregnancy obesity. Generalized estimating equations examined the associations of pre-pregnancy obesity with NNNS summary scores adjusting for maternal minority race or ethnicity, gestational diabetes, maternal hypertension (chronic or pregnancy-induced), pregnancy weight change, PMA at birth, and PMA at NNNS exam. Pre-pregnancy obesity was associated with an increase in non-optimal reflexes (B=0.41, SE=0.17, Adjusted Means: Pre-pregnancy obesity=5.64, No pre-pregnancy obesity=5.23).

Conclusions Infants of mothers who were obese prior to pregnancy showed an increase in non-optimal reflexes on the NNNS. The association between maternal pre-pregnancy obesity and atypical child neurodevelopment may be ascertained as early as infancy.

FATTY ACIDS AND GROWTH AND DEVELOPMENT IN CRITICALLY ILL NEONATES

A Sivanandam*, S Govardhan, N Li, KL Calkins. UCLA, Los Angeles, CA

10.1136/jim-2019-WMRC.55

Purpose of study Docosahexaenoic acid (DHA) and arachidonic acid (ARA) are essential for development. Infants in the neonatal intensive care unit (NICU) are at risk for ARA and DHA deficiencies secondary to premature delivery and prolonged parenteral nutrition (PN). This study’s purpose is to determine the correlation between DHA and ARA and growth and neurodevelopment.

Methods used Inclusion criteria for this prospective study 1) <14 days of age, 2) PN-dependence and 3) follow-up in the High-Risk Infant Follow-Up (FU) Clinic. Blood samples were collected at enrollment and weekly on PN. Gas chromatography/mass spectrometry was used to measure DHA and ARA in the red blood cell membrane. Neurodevelopment was assessed using the Bayley Scales of Infant Development (BSID-III).

LONG-TERM NEURODEVELOPMENTAL OUTCOMES AMONG PRETERM INFANTS EXPOSED TO GESTATIONAL DIABETES MELLITUS

1,2) Sugar*, 1,2) E Lum, 1D Poelter, 1A Katheria. 1Western University of Health Sciences, Pomona, CA; 2Sharp Mary Birch Hospital for Women and Newborns, San Diego, CA

10.1136/jim-2019-WMRC.54

Purpose of study Gestational Diabetes Mellitus (GDM) is associated with adverse pregnancy outcomes. Current evidence suggests the potential adverse effects GDM has on fetal brain development and neurodevelopmental delays. The purpose of this study is to investigate the relationship between infants born to mothers diagnosed with GDM and neurodevelopmental outcomes at two-year follow-ups.

Methods used This is a retrospective, multicenter cohort study from May 2007 to July 2019 of preterm infants. Infants were assessed using the Bayley Scales of Infant and Toddler Development (Bayley-III), which screens for cognitive, language, and motor delays in early childhood. Multivariable regression analyses were used to determine an association between GDM exposure and Bayley scores. Independent variables considered for inclusion in multivariable modeling included gestational age, maternal age, length of stay in the NICU, Apgar score at 5 minutes, days of ventilation, birth weight, intraventricular hemorrhage, retinopathy of prematurity, and bronchopulmonary dysplasia.

Summary of results 748 infants with 130 (17%) born to mothers diagnosed with GDM were assessed. Multivariable regression analysis showed a significant decrease in two-year cognitive, language, and motor composite scores when infants where exposed to GDM (unstandardized-β [95% CI]: -4.312[-7.587 to -1.036], P=0.01; -6.347 [-10.693 to -2.0], P=0.004; -3.582 [-7.084 to -0.079], P=0.045, respectively).

Conclusions This study found infants exposed to GDM during pregnancy had an increased risk of poorer cognitive, language, and motor outcomes at two-year follow-ups. Larger prospective studies are needed to confirm this association and ascertain whether neurodevelopmental delays persist from two years of age into later childhood.

ARA Changes Correlate with language development
Abstracts

56 ASSOCIATIONS BETWEEN CHRONIC LUNG DISEASE, CRY ACOUSTICS, AND NEONATAL NEUROBEHAVIOR IN INFANTS BORN BEFORE 30 WEEKS GESTATION
M Martin*, L Smith, S Sheinkopf, J Hofheimer, E McGowan, T O'Shea, S Pastyrnak, C Neal, B Carter, J Helderman, J Check, A Soliman, M Roberts, L Dansereau, S DellaGrotta, B Lester. The NOVI Research Network; Providence, RI

Purpose of study Chronic lung disease (CLD) is a known risk factor for developmental delays in infants born preterm. Most studies have focused on neurodevelopmental exams performed at or after 12 months of age, with no known studies in patients prior to discharge from the NICU. Our aim is to examine whether the presence of CLD at or near the time of hospital discharge is associated with short-term neurobehavioral outcomes.

Methods used The Neonatal Neurobehavior and Outcomes in Very Preterm Infants (NOVI) study recruited from 9 NICUs affiliated with 6 universities participating in the Vermont-Oxford Network. The NICU Network Neurobehavioral Scale (NNNS) and cry acoustics were used to assess neurobehavior at discharge. Maximum likelihood factor analysis was performed on cry acoustics. Generalized estimating equations examined the associations of CLD with NNNS summary scores and cry factors adjusting for maternal medical risks, severe ROP, brain injuries, infant sepsis, outborn status, antenatal steroid use, post menstrual age (PMA) at birth and PMA at NNNS exam.

Summary of results 709 infants enrolled in NOVI; 418 had complete medical, NNNS, and cry data and were analyzed. Of those, 197 (47.1%) infants had CLD. Factor analysis of cry data revealed two factors that explained the greatest proportion of variation: frequency/energy (loudness) and hyperphonation (high pitch). Cry factors were not associated with CLD. CLD was associated with a decrease in the NNNS attention summary score (B=-0.34, SE=0.16, Adjusted Means: CLD=4.63, No CLD=4.98).

Conclusions The presence of CLD at the time of NICU discharge was associated with a decrease in attention scores on the NNNS. CLD has been shown to have long-term adverse effects on behavior including attentional skills. Earlier identification of these at risk infants may lead to earlier initiation of therapeutic interventions.

57 PROCALCITONIN IN NEONATES UNDERGOING EARLY ONSET SEPSIS WORK UP
A Zepeda-Tiscareno*, MG Domínguez García, FB Wertheimer, A Chambilo, M Cielo, R Ramanathan, M Durand. LAC + USC Medical Center, Keck School of Medicine; University of Southern California, Los Angeles, CA

Purpose of study Obtain serum procalcitonin (PCT) reference values for neonates undergoing early onset sepsis (EOS) work up at LAC+USC Medical Center. Evaluate the utility of PCT when compared to high sensitivity C-reactive protein (hsCRP) in those who are pending blood culture results.

Methods used Demographic and laboratory values (blood culture, CBC, PCT, hsCRP) of neonates undergoing EOS work up admitted to the nursery over a 6 month period were analyzed. Data were retrospectively collected from our electronic medical records. PCT values were obtained by electrochemiluminescence immunoassay and hsCRP by immunoturbidimetric assay.

Summary of results There were 50 patients ≥35 weeks (gestational age 38.5±1.8 weeks, birth weight 3313±614 g, mean ± SD). Paired hsCRP and PCT values were obtained prior to 6 hours, 6 to 24 hours, and greater than 24 hours of life. There were 44 healthy infants (negative work up) and 6 treated with antibiotics for presumed sepsis (one positive blood culture). The median (IQR) reference values for the healthy infants are shown in table 1; their hsCRP and PCT values are correlated with each other at <6 hours (Pearson correlation coefficient, r=0.36, P=0.04), but are more highly correlated at 6–24 hours (r=0.47, P=0.0036). The hsCRP median values for the 6 infected newborns were 2.40, 17.30, and 20.60 mg/L and the PCT median values were 1.47, 24.59, and 17.89 ng/mL, respectively.

Abstract 57 Table 1 Biomarker values in healthy newborns

<table>
<thead>
<tr>
<th>Biomarker (mg/L)</th>
<th>6 to 24 hrs (n=36)</th>
<th>&gt;24 hrs (n=37)</th>
</tr>
</thead>
<tbody>
<tr>
<td>hsCRP (mg/L)</td>
<td>0.20 (0.20 – 0.40)</td>
<td>1.75 (0.83 – 4.05)</td>
</tr>
<tr>
<td>Procalcitonin (ng/mL)</td>
<td>0.17 (0.13 – 0.28)</td>
<td>1.53 (0.81 – 4.47)</td>
</tr>
<tr>
<td></td>
<td>1.38 (0.81 – 3.57)</td>
<td>1.38 (0.81 – 3.57)</td>
</tr>
</tbody>
</table>

Conclusions PCT reference values for healthy newborns with negative EOS work up are presented. PCT and hsCRP values are highly correlated at 6–24 hours and at greater than 24 hours. Studies with a larger number of healthy and sick newborns are needed, especially in very preterm infants.

58 EFFECTS OF CLINICAL AND HISTOLOGIC CHORIOAMNIONITIS ON OUTCOMES OF VERY LOW BIRTH WEIGHT PRETERM INFANTS
S Sakhamuru*, A Hisey, L Barton, R Ramanathan, M Binwale. LAC+USC Medical Center, Los Angeles, CA

Purpose of study Clinical and histologic chorioamnionitis complicate many preterm births with preterm labor or premature rupture of membranes, and has been shown to be a risk

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10.1136/jim-2019-WMRC.57

10.1136/jim-2019-WMRC.58
factor of adverse neonatal outcomes, including earlier gestational age at delivery and neonatal brain and lung injuries. The goal of this study is to evaluate clinical and histologic chorioamnionitis as risk factors for short and long term neonatal morbidities.

Methods used This is a retrospective observational study with data gathered for very low birth weight (VLBW) infants, defined as less than 1500 grams, born at LAC+USC Medical Center between 2009–2018. Clinical chorioamnionitis data was obtained based on maternal clinical symptoms diagnosed by the obstetrician. Histologic chorioamnionitis was derived from certified placenta pathology reports. Delivery resuscitation efforts and the most common neonatal morbidities and mortality were collected.

Summary of results Of the 308 infants, 95.1% had clinical chorioamnionitis and 64% had histologic chorioamnionitis. Infants with histologic chorioamnionitis had increased risk for intubation at the time of delivery (61.5% vs. 42.5%, P=0.003) and were likely to remain intubated at 24 hours of age (54.7% vs. 39.3%, P=0.020). There was also an increased risk of intraventricular hemorrhage (IVH) in this group (47.3% vs. 32.5%, P=0.021). The infants with clinical chorioamnionitis had a significant risk of having severe IVH (14.1% vs. 2.5%, P=0.001). Finally, those with histologic chorioamnionitis had a higher risk of mortality (17.6% vs. 7.7%, P=0.024).

Conclusions VLBW infants with exposure to either clinical or histologic chorioamnionitis are at a higher risk for intraventricular hemorrhage. In addition, infants with histologic chorioamnionitis require increased intubations and are at an increased risk for mortality.

PRACTICES & ATTITUDES TOWARD COMPASSIONATE EXUDBATION IN THE NEONATAL INTENSIVE CARE UNIT

1,2KE Goodman*, 3M Fossa, 3S Dabagh, 3M Caliboso, 3S Nair, 3D Lotten, 1LAC+USC Medical Center, Keck School of Medicine of USC, LA, CA; 3Fetal and Neonatal Institute, Children’s Hospital Los Angeles, Keck School of Medicine of USC, LA, CA; 3Children’s Hospital Los Angeles, Keck School of Medicine of USC, LA, CA.

Purpose of study Compassionate Exubration (CE) in the Neonatal Intensive Care Unit (NICU) may be pursued when long-term survival with acceptable quality of life is not possible, yet little medical literature exists to guide best practice. A multidisciplinary quality improvement (QI) project assessed baseline practices and attitudes among NICU providers and created an intervention to improve quality and standardization of CE in a Level 4 freestanding children’s hospital.

Methods used 12 months of retrospective analysis and chart review assessed practices, attitudes and symptom management among NICU providers towards CE. Staff who cared for a patient who underwent CE were surveyed on their comfort and team communication. A standardized CE checklist and debrief were then created to address common factors in CE management. After provider education, this intervention is being prospectively evaluated.

Summary of results In our NICU, 60% (30 of 50) of deaths in a 12 month period occurred after CE. Provider survey analysis found consistency in choices for PRN and standing pain medications, but high variability in other symptom management. Nursing survey analysis found 80% of nurses who cared for a patient who underwent CE (n=44) found the medical plan for CE adequate. 79% (n=28) found it easy to maintain the patient’s comfort. Only 12% of nurses debriefed with the medical team after CE, and 57% rated the communication with the medical team as ‘good’. Analysis after checklist and debrief implementation is ongoing, with 8 CE events pending chart review and post-implementation survey.

Conclusions While retrospective analysis found consistency regarding pain management, there was variation in other management aspects of CE. A notable minority of nurses showed concern over maintenance of patients’ comfort, and debriefings were not routine. We will use feedback from checklist and debrief testing to modify tools for NICU practice.
Neonatology pulmonary I
Concurrent session
12:45 PM
Thursday, January 23, 2020

61 POTENTIAL ROLE OF FOX FAMILY TRANSCRIPTION FACTORS IN THE PATHOGENESIS OF BRONCHOPULMONARY DYSPLASIA
A Raheed*, F Gao, S Belluscio, R Ramanathan, P Minoo. LAC+USC Medical Center and Children’s Hospital Los Angeles, Los Angeles, CA
10.1136/jim-2019-WMRC.61

Purpose of study Bronchopulmonary dysplasia (BPD) is a chronic lung disease most commonly seen in preterm infants as a result of long-term mechanical ventilation and oxygen exposure. The pathogenesis resulting in BPD is not completely understood. Studies using postmortem BPD lung tissue have shown that there is altered expression of a number of developmentally important genes.

Methods used We generated three independent genetic models of BPD-like phenotype by disrupting the signaling pathways of TGFβ, PDGFα, and IGF-1 in neonatal lung in mice. The mutations to disrupt the pathways were induced in secondary crest myofibroblasts (SCMF) whose role is critical for normal alveologenesis. RNAseq analysis was performed which identified a cluster of differentially expressed genes that were common amongst the three BPD-like phenotypes. To assess the physiologic relevance of a specific subset of the identified genes, we examined their expression in a well-established hyperoxia-induced hypoalveolization mouse model. We analyzed gene expression using a total of ten mice exposed to hyperoxia compared to eight combined controls at three different points during neonatal life. mRNA expression was assessed by quantitative RT-PCR.

Summary of results Expression of Foxq1, Foxd1, Foxc2 were decreased in the mouse BPD lungs suggesting a role for the selected genes in the pathogenesis of BPD. We further demonstrate that Foxd1 is expressed in SCMF and ablation of SCMF during alveologenesis via Foxd1-cre arrests alveolar formation leading to a BPD-like phenotype. To validate our findings in the mouse model, we show that Foxd1 is also reduced in human BPD lung tissue.

Conclusions Collectively, these observations suggest that disruption of Foxd1 and FOX family transcription factors might play a major role in the pathogenesis of BPD in preterm infants. This study was supported by NHLBI and The Hastings Foundation.

62 ANTENATAL EXOSOME TREATMENT PRESERVES LUNG STRUCTURE AND VASCULAR GROWTH IN AN EXPERIMENTAL MODEL OF BRONCHOPULMONARY DYSPLASIA DUE TO CHORIOAMNIONITIS
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1Pediatric Heart Lung Center, University of Colorado School of Medicine, Aurora, CO; 2University of Notre Dame, Notre Dame, IN; 3Boston Children’s Hospital, Harvard University, Cambridge, MA
10.1136/jim-2019-WMRC.62

Purpose of study Bronchopulmonary dysplasia (BPD), the chronic lung disease of prematurity, results in many complications that significantly alter quality of life and decrease life expectancy. Prenatal stresses such as chorioamnionitis (CA) increase the risk for BPD in infants independent of postnatal lung injury. Past studies have shown that antenatal endotoxin (ETX) exposure as a model of CA causes sustained disruption of lung alveolar and vascular growth, hallmark findings in BPD. Recent research has shown promising results for postnatal treatment of BPD with mesenchymal stem cell derived exosomes (MEx), however, the antenatal efficacy of MEx for the prevention of BPD is unknown. Thus, we hypothesized that antenatal treatment with MEx in an experimental rat model of CA will be sufficient to prevent the development of BPD.

Methods used Effects of antenatal MEx injections were studied using an established rat model of CA as induced by ETX. At E20, rat litters were treated with intra-amniotic injections of saline control, ETX only (100µL/sac), or ETX plus MEx. Pups were delivered near term (E22) via cesarean section. Morphometric studies to assess airspace growth (by mean linear intercept; MLI), vascular growth (vessel density; VD), and lung function (FlexiVent) were performed at 2 weeks postnatal age.

Summary of results When compared with controls, intra-amniotic injections of ETX reduced alveolar growth as reflected by increased MLI by 23.2% and reduced vascular growth (decreased VD) by 44.4% at 2 weeks postnatal age (p<0.05). In infant rats treated with antenatal MEx and ETX, MLI and VD were not different from control rats. Lung resistance and compliance also improved with antenatal MEx treatment in comparison with the effects of ETX alone.

Conclusions Antenatal treatment with MEx preserves lung structure and function in rats with experimental BPD induced by ETX. We speculate that early MEx treatment may be sufficient to prevent the development of BPD in premature infants, especially in the clinical setting of CA.

63 INTRANASAL AND INTRAPERITONEAL LIPOPOLYSACCHARIDE ADMINISTRATION ACTIVATES DISTINCT GENE NETWORKS IN LUNG MACROPHAGE SUBSETS
E Saji*, N Spann, O Zhengyu, L Prince, C Glass. University of California San Diego, San Diego, CA
10.1136/jim-2019-WMRC.63

Purpose of study Microbial pathogens attack the lung via the airspace or pulmonary circulation. The lung immune system contains distinct myeloid cell populations resident in the alveolar space, interstitium, and pulmonary capillary bed. How the anatomic site of the initial inflammatory signal might lead to differences in immune response is not fully understood.

Methods used We administered LPS intraperitoneally (i.p) or intranasally (i.n.) to C57BL/6J mice and examined transcriptional changes in lung macrophages and monocytes. Alveolar macrophages (AM), interstitial macrophages (IM), and inflammatory Ly6c+ monocytes (iMo) were sorted by FACS 2 h, 6 h and 22 h after LPS. Gene expression was measured in sorted cells by RNA-seq and chromatin accessibility by ATAC-seq. Informatic analysis included ingenuity
pathway analysis (IPA) and HOMER for transcription factor binding prediction.

Summary of results Following i.p. LPS, IM and iMo showed robust, largely overlapping changes in gene expression. AM showed modest changes. IPA of enriched pathways for iMo and IM showed substantial qualitative differences. After i.n. LPS, AM were most responsive, with few changes in gene expression in IM and iMo. Comparing genes induced at the time of maximal response in IM (2h, i.p. LPS) to genes in AM (6h, i.n. LPS) we found minimal overlap. Remarkably, LPS caused opposite changes in the most enriched gene expression pathways in AM and IM. For example, LPS stimulated expression of oxidative phosphorylation genes in AM but repressed them in IM. ATAC-seq analysis identified a distinct group of genes whose genomic accessibility correlated with cell-specific transcriptional responses.

Conclusions Different routes of LPS administration in vivo lead to very distinct transcriptional responses within lung myeloid populations. Remarkably, the direct responses of IM to i.p. LPS and AM to i.n. LPS are mostly divergent, in many cases leading to predictions of opposite biological outcomes. Differences in the chromatin landscapes of genes with divergent responses may lead to distinct enhancers available for binding signal dependent transcription factors, including NF-kB.

Summary of results IA s-Flt decreased RAC and PVD by 28% and 43%, respectively, and increased RVH by 32% as compared to controls ($p<0.001$). IA s-Flt increased lung resistance by 30% and decreased compliance by 30% compared to controls ($p<0.01$). IA 1,25-(OH)$_2$D$_3$ treatment in s-Flt exposed animals restored lung structure, function, and prevented RVH when compared to controls ($p=ns$).

Conclusions IA VD improved infant lung structure and function and prevented right ventricular hypertrophy after s-Flt exposure in vivo. We speculate that 1,25-(OH)$_2$D$_3$ may preserve lung growth and function through enhanced angiogenesis in experimental PE.

### Abstract 64

**ANTENATAL VITAMIN D ATTENUATES LUNG INJURY IN A PREECLAMPSIA MODEL OF BRONCHOPULMONARY DYSPLASIA**

M. Cookson*, T. Gonzalez, G. Seedorf, B. Smith, SH. Abman, E. Mandell. University of Colorado, Aurora, CO

10.1136/jim-2019-WMRC.64

**Purpose of study** Preeclampsia (PE) is a major risk factor for preterm birth and is strongly associated with the subsequent development of bronchopulmonary dysplasia (BPD), the chronic lung disease of prematurity. We have previously shown that antenatal (AN) exposure to soluble fms-like tyrosine kinase 1 (s-Flt), an endogenous VEGF antagonist that is markedly increased in maternal blood and amniotic fluid in PE, causes abnormal lung structure and function in infant rats. Clinical studies suggest that maternal vitamin D deficiency is a risk factor for severe PE, however, whether AN vitamin D (VD) treatment can restore lung structure and function after exposure to AN s-Flt is unknown. We aimed to determine if early VD (1,25-(OH)$_2$D$_3$) treatment will preserve lung structure and function in infant rats after antenatal exposure to s-Flt.

**Methods used** Fetal rats were exposed to recombinant human sFlt-1 (1μg), recombinant human s-Flt (1μg) + 1,25-(OH)$_2$D$_3$ (1ng/ml), or saline via intra-amniotic (IA) injection at E20 and delivered two days later. At 14 days of age, lung function including total respiratory system compliance (Crs) and resistance (Rrs) was determined by Flexivent and lung structure was assessed for radial alveolar counts (RAC) and pulmonary vessel density (PVD) by standard morphometric analysis. Infant hearts were assessed for right ventricle hypertrophy (RVH) by the ratio of RV/LV+S (Fulton’s Index).

**Summary of results** IA s-Flt decreased RAC and PVD by 28% and 43%, respectively, and increased RVH by 32% as compared to controls ($p<0.001$). IA s-Flt increased lung resistance by 30% and decreased compliance by 30% compared to controls ($p<0.01$). IA 1,25-(OH)$_2$D$_3$ treatment in s-Flt exposed animals restored lung structure, function, and prevented RVH when compared to controls ($p=ns$).

**Conclusions** IA VD improved infant lung structure and function and prevented right ventricular hypertrophy after s-Flt exposure in vivo. We speculate that 1,25-(OH)$_2$D$_3$ may preserve lung growth and function through enhanced angiogenesis in experimental PE.

### Abstract 65 Table 1

<table>
<thead>
<tr>
<th>Expression in VD deficiency</th>
<th>Gene</th>
</tr>
</thead>
<tbody>
<tr>
<td>Upregulated</td>
<td>ACTRI1A, LRP1, PHF5A, CELF2, EFEMP2, GLB1, IGFAP1, PRRC2A, PMM1</td>
</tr>
<tr>
<td>Downregulated</td>
<td>ARL3, SMA3D, PA2G4, NHP2L1, RASD9, I1LTR1, PHB, ATG5, AAGAB, TNAFAPI, GSAP, STAT6</td>
</tr>
</tbody>
</table>

**AAGAB, TNFAIP8, GSAP, STAT6**
from GWAS were found to have differential expression in accordance with VD status. The up- or down- regulation of these genes in VD deficient rats (no dietary D3) in comparison to their expression in VD supplemented groups is as shown in the table 1.

Conclusions Perinatal VD status determines the expression of several asthma-related genes as identified by GWAS. Functional network data mining places VD as a key determinant of several lung development pathways that determine lung homeostatic vs. asthmatic phenotypes.

**Abstract 66**

VITAMIN D STIMULATES PULMONARY ENDOTHELIAL CELL GROWTH THROUGH ENHANCED PPARγ SIGNALING

1T Gonzalez*, 1G Seedorf, 1J Fleet, 1SH Abman, 1E Mandell. 1University of Colorado Denver, Aurora, CO; 2Purdue University, West Lafayette, IN

10.1136/jim-2019-WMRC.66

Purpose of study Maternal vitamin D deficiency (VDD) in pregnancy increases risk for childhood lung disease. Past studies have shown offspring of VDD dams have impaired lung function, airway hyperreactivity, and decreased lung vascular growth at two weeks. We further found vitamin D (VD) stimulates fetal pulmonary artery endothelial cell (PAEC) growth and distal lung structure in vivo, but mechanisms through which VD enhances angiogenesis are uncertain. Peroxisome proliferator-activated receptor (PPARγ) regulates alveolar development and PPARγ expression is decreased in lungs of rats with VDD. The purpose of this study is to determine whether VD stimulates PAEC growth through enhanced PPARγ signaling or upregulation of vascular endothelial cell growth factor (VEGF), a pro-angiogenic agonist.

Methods used PAEC were isolated from proximal pulmonary arteries of late gestation fetal sheep. Three-day growth assays were performed on PAEC treated with either VEGF or VD (1,25-(OH)2 D3) in the presence or absence of T007 (a specific PPARγ antagonist), Pioglitazone (Pio; a PPARγ agonist), or Axitinib (Ax; a VEGF receptor inhibitor). PAEC lysates from these experiments were collected for assays of VEGF and PPARγ protein.

Summary of results Treatment with either VD, VEGF or Pio alone increased cell growth above basal conditions by over 30% (p<0.0001). PAEC exposed to either T007 or Ax alone did not change cell growth compared to controls. PPARγ inhibition of PAEC prevented VD and Pio enhancement of PAEC growth. Conversely, PAEC exposed to T007 and treated with VEGF demonstrated a 20% increase in cell growth (p<0.001). However, VD stimulated PAEC growth during VEGF inhibition by 20% growth compared to controls (p<0.0001). VD treatment of PAEC increased PPARγ and VEGF expression by 80% compared to controls (p<0.001).

Conclusions We conclude that the mechanism of VD treatment on PAEC growth cannot be explained by increased VEGF expression alone; this VEGF-independent component of VD induced PAEC growth is likely due to the independent effects of VD on PPARγ signaling.

**Abstract 67**

DHA SUPPLEMENTATION OF GROWTH RESTRICTED RATS INCREASES AIRWAY HYPERRESPONSIVENESS IN ASSOCIATION WITH DECREASED NUCLEAR FABP4


10.1136/jim-2019-WMRC.67

Purpose of study Preterm infants often experience postnatal growth restriction (PGR). PGR increases the severity and incidence of sex-divergent lung outcomes including bronchopulmonary dysplasia (BPD) and airway hyperresponsiveness (AHR). Docosahexaenoic acid (DHA) is important for lung development and inflammation control. A molecular mediator of DHA in the lung is fatty acid binding protein 4 (FABP4). FABP4 and DHA are decreased in BPD and in AHR. However, outcomes from clinical studies examining the effects of DHA supplementation on lung outcomes are conflicting. As a result, consensus on dose and sex effects of DHA supplementation in human PGR infants is lacking. We previously showed in a rat model that PGR causes sex-divergent 1) baseline deficits in lung function, 2) changes in whole cell FABP4 protein, and 3) altered circulating DHA. We hypothesize that in PGR rats, postnatal DHA supplementation causes dose and sex-dependent changes in response to methacholine challenge in association with changes in nuclear FABP4.

Methods used We induced PGR by randomizing newborn rat pups into litters of 8 (control) or litters of 16 (PGR). Each litter was randomized to receive diets with DHA at 0%, 0.01%, or 0.1%. At d24 of life, methacholine challenge was performed using the FlexiVent. Lung nuclear and cytoplasmic FABP4 protein levels were determined using western blot.

Summary of results Rat pups in the PGR group weighed less than control through d21 on all DHA diets. In male PGR rats, DHA at 0.1% worsened response to methacholine (increased airway resistance and decreased lung compliance) relative to male control and male PGR rats not receiving DHA. In addition, nuclear FABP4 was reduced in male PGR rats receiving the 0.1% DHA diet. Female PGR rats with or without DHA did not differ in response to methacholine or in nuclear FABP4.

Conclusions We conclude that in PGR rats, postnatal DHA supplementation causes dose and sex-dependent changes in response to methacholine challenge in association with
Use of full-coverage dressing with closed incision negative pressure therapy for bilateral mastectomy in breast cancer patients

Purpose of study Bilateral mastectomy with immediate reconstruction, a complex surgery undertaken to prevent or treat breast cancer, results in incisions at risk for post-surgical complications. Incisions created during mastectomy often face challenges with wound healing due to well-documented negative impacts of certain cancer therapies. These incisions may benefit from closed incision negative pressure therapy (ciNPT)—a wound dressing that holds incisions together, facilitates removal of fluid and infectious materials, and creates a barrier to external contaminants, supporting tissue healing and repair. A two-piece, full-coverage dressing designed to give negative pressure to incisions and surrounding tissues is available for use with commercial ciNPT systems. This case series reports our initial experience using this dressing with ciNPT in 3 breast cancer patients who underwent bilateral mastectomies with immediate reconstruction.

Methods used Full-coverage dressings were placed over the breast once bilateral mastectomy, reconstruction, and closure was complete. ciNPT was initiated on postoperative day (POD) 0 and left in place for 5–6 days. Oral antibiotics were administered post-surgery per best practice.

Summary of results The 3 patients in our case series underwent bilateral mastectomy with immediate reconstruction. Patient 1, a 59-year-old female with type 2 diabetes, hypertension, and obesity, had bilateral vertical incisions. Patient 2 and 3 were 58- and 50-year-old females with notable medical histories. Patient 2 had bilateral inverted ‘T’ incisions. Patient 3 had bilateral inframammary incisions. After application of ciNPT, all patients were discharged POD 1 and returned POD 5–6 to remove dressings. Follow-up at POD 30 showed closed incisions in all patients. There were no incidences of seromas, dehiscences, or site infections. At follow-up 4–5 weeks post-surgery no adverse events were reported.

Conclusions In this patient group at risk for poor wound healing due to cancer therapy, full-coverage dressings with ciNPT supported incision healing and soft tissue repair after bilateral mastectomy and reconstruction for breast cancer.
of thyroidectomies due to indeterminate nodules by FNAB alone. Therefore, there is a need for a more accurate and time-efficient diagnostic approach for analyzing indeterminate thyroid nodules. Recently, the osteogenic protein Enigma has been associated with different cancer types, including thyroid cancer progression and calcification through its interaction with bone morphogenic protein-1 (BMP-1), and tyrosine kinases linked to mitogenic signaling pathways. Our published data on Enigma protein analysis with immunohistochemistry showed promising results in discriminating between malignant versus benign thyroid nodules, and demonstrated correlation with thyroid cancer staging. In this study, we are investigating Enigma at a gene expression level by RT-qPCR, which is a quantitative and more time-efficient method that requires smaller samples (FNA) than immunohistochemistry.

Methods used We extracted mRNA/DNA/proteins from fresh malignant and benign thyroid nodules using a Qiagen DNA/RNA/Protein Kit. We prepared cDNA from isolated pure mRNA and ran through Enigma-qPCR assay using primers.

Summary of results The results showed that the Enigma-mRNA expression level was 3-fold higher in malignant as compared to benign thyroid tissue, which is statistically significant. This finding supports our previous Enigma immunohistochemistry data and shows a relative quantitative difference in Enigma-mRNA expression level between malignant and benign thyroid nodules.

Conclusions We conclude that Enigma-RT-qPCR can be used to effectively determine malignancies in FNAB samples derived from thyroid nodules. This method could potentially enhance the diagnostic accuracy of indeterminate nodules and decrease diagnostic thyroidectomies and subsequent morbidity.

71 PERIPAPILARY RETINAL NERVE FIBER LAYER MICROCIRCULATION IN GLAUCOMA PATIENTS OF AFRICAN AND EUROPEAN DESCENT USING OPTICAL COHERENCE TOMOGRAPHY ANGIOGRAPHY


10.1136/jim-2019-WMRC.71

Purpose of study To investigate the differences in peripapillary retinal nerve fiber layer (RNFL) microcirculation in open angle glaucoma (OAG) patients of African descent (AD) and European descent (ED) using optical coherence tomography-angiography (OCTA).

Methods used Twenty-three OAG eyes of AD and 27 OAG eyes of ED were scanned using AngioPlex OCTA system (Zeiss Meditec Inc., Dublin, CA) covering a 6 × 6 mm scanning area centered at the optic nerve head (ONH). Peripapillary RNFL microcirculation was measured by calculating the overall blood flux (BF) and vessel area density (VAD) within an annulus centered at the ONH (2.5-mm and 3.7-mm as inner and outer diameters) with the exclusion of big retinal vessels. Two-sample, independent t-tests were used to compare the peripapillary RNFL blood flow metrics between AD and ED eyes. Linear regression models were used to investigate the correlation between peripapillary RNFL blood flow metrics and structural and functional parameters.

Summary of results There was no significant difference in age, sex, systolic blood pressure (BP), diastolic BP, mean ocular perfusion pressure, RNFL thickness, visual field (VF) mean deviation (MD) and VF pattern standard deviation (PSD) between AD and ED glaucomatous eyes included (p ≥ 0.062). Eyes of AD and ED had similar peripapillary RNFL blood flow metrics (p≥0.380). Peripapillary RNFL blood flow metrics were significantly correlated with VF MD, VF PSD and RNFL thickness (p=0.026).

Conclusions Peripapillary RNFL blood flow metrics were significantly correlated with disease severity in AD and ED glaucomatous eyes. Peripapillary RNFL microcirculation detected by OCTA was similar between AD and ED glaucomatous eyes.

TISSUE PLASMINOGEN ACTIVATOR PROTOCOL FOR FROSTBITE IN ALASKA: IMPROVED ACCESS IN RURAL AND REMOTE AREAS

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10.1136/jim-2019-WMRC.72

Purpose of study Use of tissue plasminogen activator (tPA) for treatment of severe frostbite has facilitated limb salvage when previously amputation was inevitable. To address Alaska’s rural and remote areas, a protocol for systemic tPA in patients with grade 3 or 4 frostbite extremity injuries was adopted. This study evaluated the efficacy of this protocol and identified barriers to its success.

Methods used We compiled all patients treated for frostbite from this institution from October 2012 – May 2019 using data from inpatient admissions, outpatient clinic visits, Emergency Department consult notes, and pharmacy records of tPA infusions. This established a pre-treatment cohort from October 2012 – September 2017 and a tPA eligible cohort from October 2017 – May 2019. Chart review determined key variables including injury grade, timing and location (of injury, first evaluation, and tPA administration), distance from treating facility, amputation level, and tPA candidacy. Descriptive statistics were gathered, and chi squared test was used to compare the two groups.

Summary of results Clinical characteristics were similar in both groups. A pre-protocol group who were retrospectively determined to be tPA candidates consisted of 13 men (76%) with an average age of 36.3 [25 (26–51)]. Post-protocol tPA recipients consisted of 7 men (78%) with an average age of 37.2 [20 (25–45)]. Amputation of grade 3 and 4 frostbite injuries decreased from 82% (pre-treatment) to 22% (post-treatment). Average time from rewarming to tPA administration was 5.1 hours [3.9 (3.4–7.3)]. There were no adverse events of tPA treatment. Of the two amputations after tPA, one was complicated by infection, and one received amputation after a second re-freezing injury. Many patients were not considered for tPA based on time from rewarming to evaluation.

<table>
<thead>
<tr>
<th>Abstract 72 Table 1</th>
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<tbody>
<tr>
<td><strong>Frostbite Cases</strong></td>
</tr>
<tr>
<td>Pre-protocol (2012–17)</td>
</tr>
<tr>
<td>Post-protocol (2017–19)</td>
</tr>
</tbody>
</table>

A30 JIM 2020;68(Suppl 1):A1–A228
Conclusions Systemic tPA administration is shown to be effective in decreasing amputation among patients with grade 3 and 4 frostbite injuries while providing means of accessible treatment in rural and remote settings. Education of frostbite as a time-critical injury may increase patient identification for treatment and improve outcomes.

**NECROTIZING ENTEROCOLITIS IN A RAT ANIMAL MODEL**

1. GY Perrier*, 2. YS Mendez, 3. FA Khan, 4. A Radulescu. 1. Loma Linda University, Loma Linda, CA; 2. Loma Linda University Health, Loma Linda, CA

10.1136/jim-2019-WMRC.73

Purpose of study Necrotizing Enterocolitis (NEC) is one of the leading causes of death in premature infants. The purpose of this research endeavor was to reproduce and develop an animal model of NEC that can be used for possible therapeutic interventions.

Methods used Sprague Dawley rat pups were delivered prematurely by C-section at 21 days gestation. Newborn rat pups were feed by an orogastric tube with 15 g Similac & 75 cc of Esbilac 200 kcal/kg every 4 h, receiving 0.1 ml on day 1 of life advancing to 0.4 ml by day 4 of life. Pups were exposed to 100% N x 60 sec followed by 40C x 10 min beginning 10 min after birth three times a day. The pups received LPS (2 mg/kg) via OG tube 8h after birth. Pups were monitored closely for clinical signs of NEC (bloody stools, abdominal distention, lethargy, respiratory distress) and were sacrificed if any above symptoms were observed. On day 5, all surviving pups were sacrificed and intestines were harvested. Samples of duodenum, jejunum, ileum and colon were fixed in 10% formalin. Histological changes in the intestines were graded: grade 0, normal, no damage; grade 1, epithelial cell lifting or separation; grade 2, sloughing of epithelial cells to mid villus level; grade 3, necrosis of entire villus; and grade 4, transmural necrosis. Tissues with histological scores 2 or higher were designated as positive for NEC.

Summary of results The incidence of NEC found in our experiment using preterm rat pups was 62.9%.

Conclusions The rat animal model of NEC is a useful tool in the study of NEC as well as in developing therapeutic interventions. Currently, our laboratory is investigating the role of Heparin-binding EGF-like growth factor (HB-EGF) in prevention of NEC using this animal model.

**SURGICAL CAPACITY AND TRAUMA SYSTEM FUNCTIONALITY IN RURAL UGANDA**


10.1136/jim-2019-WMRC.74

Purpose of study Surgical capacity assessment in combination with an evaluation of the trauma system and services at Soroti Regional Referral Hospital (SRRH), Soroti, Uganda, are key factors to identifying priorities in care and growth in partnership together with BC Children’s Hospital. As such, study objectives include assessing surgical capacity at SRRH and evaluating functionality of the Soroti trauma system and services.

Methods used The Global Assessment in Pediatric Surgery (GAPS) checklist, a novel capacity assessment tool, was implemented at SRRH, administered via interview with a senior surgical faculty member. To evaluate the trauma system, a 26-item environmental scan was implemented via three structured interviews with a general surgeon, medical officer, and intern, in addition to a focus group with four interns. This project took place during July 2019.

Summary of results GAPS highlights strengths in surgical capacity at SRRH, including ability to perform surgery 24-hr/day; consistent access to 2 equipped operating rooms; and availability of a surgical and anesthesia team with experience in pediatric care. Over 50% of health care practitioners involved in pediatric care attend 1–2 CME sessions annually. Areas for growth in capacity include improving timely access to surgical care (currently, <50% of patients reach definitive care in <2 hrs) and formalized tracking of patient outcomes. Key identifications of the environmental scan include the need for establishment of EMS in the community, and significant trainee interest (consensus amongst 2/2 trainee interviewees, and 4/4 trainees in focus group) in the development of trauma care provider education during internship, such as ATLS and simulation training.

Conclusions This study demonstrates insight into the current scope of surgical capacity and trauma services at SRRH. Potential partnership priorities include a focus on integrated pre-hospital care and educational resources for trainees at the institution.

**COMPARISON OF SVF VIABILITY FROM FAT HARVESTED VIA ULTRASOUND ASSISTED LIPOSUCTION VERSUS STANDARD ASSISTED LIPOSUCTION: A PRELIMINARY VOLUMETRIC ANALYSIS**

1. L Tobing*, C Lee, S Roddick, CS Drew, S Gupta. Loma Linda University, Loma Linda, CA

10.1136/jim-2019-WMRC.75

Purpose of study Interest in the stromal vascular fraction (SVF) of adipose tissue has grown because of its potential to be a source of therapeutic progenitor cells due to its multipotency and ease of harvest via liposuction. The gold standard method of suction-assisted liposuction (SAL) has been established but alternative methods are gaining popularity such as ultrasound-assisted liposuction (UAL). Studies have not yet directly compared these two methods regarding the quantity of adipose-derived mesenchymal stem cells preserved. This study evaluated if ultrasound collection of fat allows for larger amounts...
of viable stromal vascular fraction after the isolation process compared to standard suction technique of fat collection in patients undergoing autologous fat transfer.

Methods used Autologous lipoaspirates were harvested from a total of eight female patients. In our study, our UAL was specifically Vibration Amplification of Sound Energy at Resonance (VASER). Lipoaspirates were cooled and stored in a freezer until processing, centrifuged in 10 mL tubes at 3000 revolutions per minute. Then, SVF was preserved by extracting and discarding the fat and plasma layer and measured for viability using trypan blue solution staining and microscopic high power field counts to determine the number of cells preserved from each method.

Summary of results Table 1 summarizes the findings of this experiment. The average mass of SVF harvested via SAL was 0.75 g versus an average of 1.64 g for the SVF harvested via UAL. A two tailed t-test revealed a p value of <0.002. Viable cell counts per high power field yielded a value of 23.2 for SAL versus 40.2 for UAL. The p value for the two tailed t-test was 0.03.

Conclusions The fat collected by the VASER method resulted in a higher volume of SVF with a greater density of viable cells. This suggests that UAL is a more efficient method of collection which also was reported to be less physically taxing for surgeons.

Abstract 75 Table 1: Summarizes the findings of this experiment

<table>
<thead>
<tr>
<th>Patient</th>
<th>Age</th>
<th>Harvest Site</th>
<th>Method</th>
<th>SVF Mass (per 10 ml)</th>
<th>Viable cells per HPF</th>
</tr>
</thead>
<tbody>
<tr>
<td>1</td>
<td>52</td>
<td>abdomen</td>
<td>SAL</td>
<td>0.62 g</td>
<td>16</td>
</tr>
<tr>
<td>2</td>
<td>49</td>
<td>outer thigh</td>
<td>SAL</td>
<td>0.72 g</td>
<td>22</td>
</tr>
<tr>
<td>3</td>
<td>50</td>
<td>abdomen</td>
<td>UAL</td>
<td>1.59 g</td>
<td>32</td>
</tr>
<tr>
<td>4</td>
<td>51</td>
<td>abdomen</td>
<td>UAL</td>
<td>0.75 g</td>
<td>33</td>
</tr>
<tr>
<td>5</td>
<td>48</td>
<td>flank</td>
<td>UAL</td>
<td>1.49 g</td>
<td>44</td>
</tr>
<tr>
<td>6</td>
<td>48</td>
<td>abdomen</td>
<td>UAL</td>
<td>1.64 g</td>
<td>38</td>
</tr>
<tr>
<td>7</td>
<td>54</td>
<td>abdomen</td>
<td>SAL</td>
<td>3.19 g</td>
<td>51</td>
</tr>
<tr>
<td>8</td>
<td>51</td>
<td>abdomen</td>
<td>UAL</td>
<td>1.45 g</td>
<td>30</td>
</tr>
</tbody>
</table>

Purpose of study Increasing organ scarcity has motivated transplant centers to relax restrictions to donation creating the term “extended-criteria donor” (ECD). ECD include organs that carry increased risk of impaired allograft function and/or donor-transmitted disease. Due to a better understanding in the utilization of these grafts, we hypothesize that both patient and graft survival associated with utilization of liver grafts from ECD will be similar to standard criteria donors (SCD).

Methods used Retrospective cohort analysis of adult patients who received a liver transplant at the University of Washington (UW) between January 1, 2014 and December 31, 2016. Clinical data was obtained from transplant database at UW. Analysis was limited to transplant recipients who received primary liver transplant. Patients with re-transplantation, multi-organ transplants and living donor liver recipients were excluded. Demographic data was analyzed using a t-test for continuous variables and a Fisher’s Exact test for categorical variables. Survival curves were calculated with Kaplan-Meier analyses and compared with a log-rank test. Cox proportional hazards model was performed to determine the contribution of the recipient and donor variables on allograft and patient survival.

Summary of results We identified 104 ECD and 135 SCD recipients. Mean age for ECD donor is 37.6±14.2 years old and SCD donor is 32.5±13.5 years old (p=0.002). ECD liv- ers have shorter cold ischemia time (7.1±2.0 hours vs. 8.0±2.5; p=0.003). The rate of primary non-function, re-transplantation, length of hospital stay and liver enzymes post-op are not different between groups. Kaplan Meier patient and graft survival show no difference between recipient groups. Cox proportional hazards model showed no statistical significance between graft or patient outcomes.

Conclusions The utilization of ECD in liver transplantation does not have any impact in both patient and graft survival. These findings suggest that ECD should be considered more in an era of scarce organ availability and high wait list mortality.

Cardiovascular II
Concurrent session
3:15 PM
Thursday, January 23, 2020

77 USE OF HEPATITIS B DONORS IN HEART TRANSPLANTATION: ARE THERE CONSEQUENCES?

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Purpose of study In the past, Hepatitis B donors have been declined in patients who have not been vaccinated by the Hepatitis B vaccine. There is concern that these donors would transmit Hepatitis B to the recipient if they are not vaccinated. There is treatment available for these Hepatitis B infections, however, it has not been established as to its efficacy in patients on immunosuppression. In addition, hepatitis viruses have been reported to affect the endothelium of vital organs. Hepatitis B may injure the endothelium of the coronary vascular tree and could potentially result in a greater development of cardiac allograft vasculopathy (CAV) after heart transplantation. Therefore, we sought to assess this possibility by examining our patients who received Hepatitis B donors.

Methods used Between 2010 and 2016 we assessed 24 heart transplant patients who received a Hepatitis B donor. All of the recipients had received a Hepatitis B vaccine prior to transplantation. Endpoints include 3-year survival, 3-year freedom from CAV, 3-year freedom from non-fatal major adverse cardiac events (NF-MACE), and freedom from first year rejection, including any treated rejection, acute cellular rejection (ACR), and antibody-mediated rejection (AMR). These study patients were compared with 600 [K11] [SA2] [SA3] patients without Hepatitis B donors in a contemporary era.
Abstract 77 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>HTx Patients with a Hepatitis B donor (n=24)</th>
<th>HTx Patients without a Hepatitis B donor (n=600)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>3-Year Survival</td>
<td>87.5%</td>
<td>86.7%</td>
<td>0.941</td>
</tr>
<tr>
<td>3-Year Freedom from CAV</td>
<td>79.2%</td>
<td>88.2%</td>
<td>0.111</td>
</tr>
<tr>
<td>3-Year Freedom from NF-MACE</td>
<td>79.2%</td>
<td>81.8%</td>
<td>0.699</td>
</tr>
<tr>
<td>1-Year Freedom from ATR</td>
<td>87.5%</td>
<td>85.2%</td>
<td>0.863</td>
</tr>
<tr>
<td>1-Year Freedom from ACR</td>
<td>95.8%</td>
<td>93.3%</td>
<td>0.695</td>
</tr>
<tr>
<td>1-Year Freedom from AMR</td>
<td>95.8%</td>
<td>94.7%</td>
<td>0.847</td>
</tr>
</tbody>
</table>

Summary of results 3-year survival, freedom from NF-MACE, and freedom from first year rejection were similar between the Hepatitis B and Control groups. There is numerically a lower incidence of freedom from CAV in the Hepatitis B group compared to the control, but this was not statistically significant.

Conclusions Hepatitis B donors appear to have acceptable outcome compared to non-Hepatitis B donors after heart transplantation. Larger numbers of Hepatitis B donors will be needed to confirm these findings.

78 FAMILY ATTITUDES TOWARD GENOMIC SEQUENCING IN CHILDREN WITH CARDIAC DISEASE

1Stanford Center for Biomedical Ethics, Palo Alto, CA; 2Stanford Medicine, Palo Alto, CA; 3Stanford Center for Biomedical Ethics, Palo Alto, CA

10.1136/jim-2019-WMRC.78

Purpose of study Care for children with cardiac disease often involves difficult decisions and clinical uncertainty. Genomic Sequencing (GS) promises to improve clinical prognostics and could impact how difficult decisions are made. We sought to determine how GS results might alter family attitudes towards bedside care choices.

Methods used We conducted semi-structured interviews of 35 families at a high-volume pediatric heart center. We discussed previous experience with and understanding of GS, perceptions toward GS in real and hypothetical scenarios, and support needed for implementing GS in clinical care. Responses were analyzed using grounded theory and a codebook was developed. Researchers discussed interpretation of codes and identified and described emerging themes. Interrater reliability was 0.91.

Summary of results Three themes emerged: 1) Is knowledge beneficial? Families saw benefits in GS when allocating resources, even if it meant limitation of treatment options for their child. 2) Is giving GS results to the healthcare system safe? All families indicated mistrust of at least one facet of the medical system including insurance companies, maintenance of confidentiality, and the incentive structure in healthcare. Some also expressed distrust of direct to consumer GS testing.

Conclusions In families of children with cardiac disease, trust is lacking in perceptions of the clinical utility of GS results, in using GS results to inform difficult decisions and in clinical protections for privacy or handling of GS results. Further efforts to improve the trustworthiness of clinical GS are needed to engage family support in implementation of GS to clinical care.

ENGINEERED CARDIAC GRAFT REDUCES HEART FAILURE-MEDIATED VENTRICULAR TACHYCARDIA


Purpose of study Patients with Heart Failure with reduced Ejection Fraction (HFrEF) have a high incidence of ventricular tachycardia (VT). Though pharmacologic and invasive therapies exist for HFrEF-mediated VT, none restore the electrical stability of the infarcted myocardium. Fibroblasts are the principle component of cardiac stroma and we hypothesize that they can be utilized surgically to modulate susceptibility to ventricular tachyarrhythmia.

Methods used We performed a proof-of-concept study by surgically implanting engineered cardiac grafts epicardially in a rodent model of chronic ischemic heart failure (HF). A single graft formulation was utilized as the experimental therapy: human dermal fibroblasts cultured into a biosorbable polyglactin-910 woven scaffold. The specific aims of this study were to 1) compare susceptibility to induced VT via programmed electrical stimulation in SHAM, HF, and HF+graft rats and 2) to compare two-dimensional epicardial electroanatomic map findings between the three groups.

Summary of results HF rats (n=29) exhibited no change (1.0 ±0.1 versus 0.7±0.2, p=0.1430) in capture threshold, a prolongation (68±3 versus 52±4, p=0.0191) of effective refractory period (ERP), and an increase (72±0% versus 0%, p=0.0394) in the incidence of inducible VT compared to SHAM rats (n=29). Graft-treated rats (n=12) exhibited no change (0.8±0.1 versus 1.0±0.1, p=0.3220) in capture threshold, a prolongation (88±7 versus 68±3, p=0.0029) of ERP and a non-significant decrease (42± versus 72%, p=0.08) in the incidence of inducible VT compared to HF rats. Epicardial electroanatomic mapping revealed lower (p<0.05) monophasic action potential amplitudes in the left ventricle of HF and graft-treated HF rats compared to SHAM rats.

Conclusions An engineered cardiac graft composed of human dermal fibroblasts on a scaffold was implanted onto the epicardium of rodents with HFrEF and altered the effective refractory period of the infarcted left ventricle. This graft may also influence susceptibility to HFrEF-mediated VT via impeding the successful conversion of R-on-T phenomena to ventricular tachyarrhythmia. This proof-of-concept study confirmed the model and methods necessary to evaluate the therapeutic potential of this cell-based therapy and other novel cell-based therapies for HFrEF.
Does the Type of Explosive Brain-Death Correlate with Outcome After Heart Transplantation?

T Hage*, K Nishihara, A Shen, R Levine, M Hamilton, J Kobashigawa. Cedars-Sinai Medical Center, Los Angeles, CA

10.1136/jim-2019-WMRC.80

Purpose of study In heart transplantation, the origin of brain death may have an impact on outcome after heart transplantation. It has been reported that explosive brain death, including head trauma, gunshot to the head, and subarachnoid hemorrhage (SAH), may lead to increased catecholamine surge with damage to the donor heart as well as upregulation of inflammatory markers. Other forms of brain death such as hanging, drowning, and drug overdose may also impact outcome, but this has not been firmly established. Therefore, we sought to evaluate explosive brain death and its relationship to outcomes in the current era.

Methods used Between 2010 and 2018, we assessed 823 heart transplant patients and divided them into two groups based on whether the heart donor underwent explosive or non-explosive brain death. Endpoints included 1-year survival, freedom from the development of cardiac allograft vasculopathy (CAV, as defined by stenosis ≥30% by angiography), non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, new congestive heart failure, percutaneous coronary intervention, implantable cardioverter defibrillator/pacemaker implant, stroke), acute cellular rejection (ACR), antibody-mediated rejection (AMR), donor specific antibodies (DSA), and severe primary graft dysfunction (PGD).

Summary of results Patients from both explosive and non-explosive brain death mechanisms have comparable outcomes in terms of survival and freedom from severe PGD, CAV, NF-MACE, and rejection.

Conclusions Explosive brain death does not appear to be associated with less than optimal outcomes after heart transplantation.

Pre-Transplant Collagen Vascular Disease is Associated with an Increase in Cardiac Allograft Vasculopathy After Heart Transplantation

G Harris*, K Nishihara, A Shen, R Levine, M Hamilton, J Kobashigawa. Smidt Heart Institute at Cedars-Sinai, Los Angeles, CA

10.1136/jim-2019-WMRC.81

Purpose of study Collagen vascular disease, including disease states such as systemic lupus erythematosus, rheumatoid arthritis, and scleroderma, have auto-antibodies in common as their etiology. These patients (pts) develop an inflammatory disease state and are known to be triggered by IgG and IgM antibodies against self. It is not clear whether pts with underlying collagen vascular disease face an increased risk of developing donor specific antibodies (DSA), rejection, and cardiac allograft vasculopathy (CAV).

Methods used Between 2010-2018, we assessed 22 heart transplant pts who had pre-transplant diagnoses of collagen vascular disease specified as lupus (n=7), rheumatoid arthritis (n=9), scleroderma (n=3), and mixed connective tissue disease (n=3). Pre-transplant immunotherapy, first-year post-transplant survival, and freedom from CAV (as defined by stenosis ≥30% by angiography), non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, new congestive heart failure, percutaneous coronary intervention, implantable cardioverter defibrillator/pacemaker implant, stroke), acute cellular rejection (ACR), antibody-mediated rejection (AMR), DSA, and left ventricular dysfunction (as defined by left ventricular ejection fraction <40%) were recorded.

Summary of results Pts with pre-transplant underlying collagen vascular disease have significantly lower freedom from CAV. First year freedom from NF-MACE, rejection, and DSA appear to be similar between study and control groups. Collagen vascular disease pts who were treated with pre-transplant disease-modifying agents were then compared as a subgroup to those pts who did not have a disease-modifying agent.

Conclusions Pts with pre-transplant collagen vascular disease appear to have increased risk of CAV. Heightened immunosuppression may be warranted in this group of patients undergoing heart transplantation.
PGD-LV when severe is noted to have increased mortality. PGD-RV has been described as hemodynamics with RAP >15 mmHg, PCWP <15 mmHg, CI <2.0 L/min/m^2, or need for RV temporary support. PGD-RV has not been well studied.

**Methods used** Between 2013 and 2018, we assessed 13 HTx pts who developed PGD-RV within 24 hours after HTx and compared them to pts without PGD. Endpoints included 1-year survival, need for temporary kidney dialysis, need for RV unloading with sildenafil, first-year any treated rejection (ATR), 1-year freedom from cardiac allograft vasculopathy (CAV, as defined by stenosis ≥30% by angiography), and 1-year freedom from non-fatal major adverse cardiac events (NF-MACE, defined as myocardial infarction, percutaneous coronary intervention/angioplasty, new congestive heart failure, pacemaker/implantable cardioverter-defibrillator placement, and stroke).

**Summary of results** Pts with PGD-RV appear to have comparable outcomes to pts without PGD for survival, freedom from CAV, freedom from NF-MACE, and rejection. Pts with PGD-RV required RV unloading therapy which may have affected outcome. There was a trend for decrease in the freedom from temporary kidney dialysis in the PGD-RV group.

**Conclusions** Right ventricular primary graft dysfunction after HTx appears to have comparable outcome with good recovery. RV unloading therapy may be helpful. However, there was an associated increased need for temporary kidney dialysis within the first year after HTx. Larger numbers are warranted to confirm these findings.

### Abstract 82 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>HTx Patients with RV-PGD (n=13)</th>
<th>HTx Patients without RV-PGD (n=407)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-Year Survival</td>
<td>92.3%</td>
<td>92.9%</td>
<td>0.824</td>
</tr>
<tr>
<td>1-Year Freedom from CAV</td>
<td>100.0%</td>
<td>96.8%</td>
<td>0.532</td>
</tr>
<tr>
<td>1-Year Freedom from NF-MACE</td>
<td>84.6%</td>
<td>88.7%</td>
<td>0.543</td>
</tr>
<tr>
<td>1-Year Freedom from ATR</td>
<td>92.3%</td>
<td>88.7%</td>
<td>0.780</td>
</tr>
<tr>
<td>1-Year Freedom from RV</td>
<td>46.2%</td>
<td>90.9%</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Unloading</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>1-Year Freedom from RV</td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>Temporary Kidney Dialysis</td>
<td>76.9%</td>
<td>91.6%</td>
<td>0.053</td>
</tr>
</tbody>
</table>

**Abstract 83 Table 1**

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>Abnormal Cardiac MRI findings of myocardial edema or fibrosis (n=12)</th>
<th>Normal Cardiac MRI findings (n=14)</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-Year Subsequent Survival</td>
<td>75.0%</td>
<td>92.9%</td>
</tr>
<tr>
<td>1-Year Subsequent Freedom from CAV</td>
<td>100.0%</td>
<td>92.9%</td>
</tr>
<tr>
<td>1-Year Subsequent Freedom from NF-MACE</td>
<td>75.0%</td>
<td>78.6%</td>
</tr>
<tr>
<td>LV dysfunction with LVEF</td>
<td>75.0%</td>
<td>21.4%</td>
</tr>
</tbody>
</table>

**Purpose of study** Heart transplant patients are known to develop cardiac rejection and cardiac allograft vasculopathy (CAV) after heart surgery. The severity of these rejection episodes and the detection of rejection has not been adequate with the endomyocardial biopsy. In fact, biopsy negative rejection is seen in approximately 5% of this patient population. The use of cardiac MRI has been increasing to demonstrate myocardial edema or fibrosis. However, it has not been established what these cardiac MRI abnormalities represent and whether they can predict the development of CAV by angiography or clinical outcomes.

**Methods used** Between 2011 and 2018, we assessed 26 heart transplant patients who underwent clinically indicated cardiac MRI imaging. In most cases, cardiac function was decreased or endomyocardial biopsy was either negative or ambiguous for rejection. Patients with abnormal cardiac MRI findings of myocardial edema or fibrosis were compared to patients without findings of myocardial edema or fibrosis. These patients were followed for 1 year after the imaging study was performed. Patients from each group were assessed for 1-year subsequent (from MRI scan) survival, 1-year subsequent freedom from CAV, 1-year subsequent freedom from NF-MACE, and the presence of left ventricular dysfunction defined as left ventricular ejection fraction (LVEF) <40%.

**Summary of results** Although the numbers are small (no p-values done), there is a numerically lower survival in patients with myocardial edema or fibrosis on cardiac MRI. Cardiac dysfunction with LVEF <40% was more prevalent in the abnormal cardiac MRI group compared to the control.

**Conclusions** Cardiac MRI findings of myocardial edema or fibrosis appear to be associated with lower outcome compared to controls. Larger studies are needed to confirm the use of cardiac MRI in this patient population.
non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, percutaneous coronary intervention/angioplasty, new congestive heart failure, pacemaker/implantable cardioverter-defibrillator placement, and stroke), and 1-year freedom from rejection (any treated rejection (ATR), acute cellular rejection (ACR), antibody-mediated rejection (AMR)).

Summary of results Patients treated with <500 mg MMF/day and >2000 mg MMF/day had a trend toward decreased freedom from ATR. There was no difference in outcome between all groups in terms of 1-year survival, freedom from CAV, freedom from NF-MACE, and freedom from ACR and AMR.

Conclusions The average daily dose of MMF in the first year after HTx does not appear to have an impact on outcome. Longer follow-up will be necessary to assess whether MMF dosing correlates to long-term complications such as CAV.

### Abstract 84 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>HTx Patients</th>
<th>HTx Patients with 500–1000 mg MMF/day (n=102)</th>
<th>HTx Patients with 1001–1500 mg MMF/day (n=50)</th>
<th>HTx Patients with 1501–2000 mg MMF/day (n=46)</th>
<th>HTx Patients with &gt;2000 mg MMF/day (n=29)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>1-Year Survival</td>
<td>92.3%</td>
<td>91.2%</td>
<td>100.0%</td>
<td>93.5%</td>
<td>89.7%</td>
<td>0.249</td>
</tr>
<tr>
<td>1-Year Freedom from CAV</td>
<td>100.0%</td>
<td>95.1%</td>
<td>92.0%</td>
<td>91.3%</td>
<td>100.0%</td>
<td>0.448</td>
</tr>
<tr>
<td>1-Year Freedom from NF-MACE</td>
<td>92.3%</td>
<td>84.3%</td>
<td>84.0%</td>
<td>82.6%</td>
<td>72.4%</td>
<td>0.353</td>
</tr>
<tr>
<td>1-Year Freedom from ATR</td>
<td>69.2%</td>
<td>86.3%</td>
<td>90.0%</td>
<td>78.3%</td>
<td>72.4%</td>
<td>0.058</td>
</tr>
<tr>
<td>1-Year Freedom from ACR</td>
<td>84.6%</td>
<td>94.1%</td>
<td>96.0%</td>
<td>93.5%</td>
<td>89.7%</td>
<td>0.543</td>
</tr>
<tr>
<td>1-Year Freedom from AMR</td>
<td>92.3%</td>
<td>97.1%</td>
<td>96.0%</td>
<td>91.3%</td>
<td>93.1%</td>
<td>0.572</td>
</tr>
</tbody>
</table>

Endpoints included the following 5-year outcomes subsequent to the development of DSA: survival, freedom from CAV, freedom from non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, percutaneous coronary intervention/angioplasty, new congestive heart failure, pacemaker/implantable cardioverter-defibrillator placement, and stroke), and freedom from rejection (any treated rejection (ATR), acute cellular rejection (ACR), antibody-mediated rejection (AMR)).

Summary of results Patients with C1q+ DSA had decreased survival and freedom from NF-MACE in the 5-years following the development of DSA. These patients also had a numerical lower freedom from CAV.

Conclusions The development of C1q+ DSA appears to be correlated with poor clinical outcome. It is not clear if treating C1q+ DSA with desensitization therapy can improve outcomes.

### Abstract 85 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>HTx Patients with C1q+ DSA (n=21)</th>
<th>HTx Patients with C1q- DSA (n=29)</th>
<th>P-Value</th>
</tr>
</thead>
<tbody>
<tr>
<td>5-Year Subsequent Survival</td>
<td>52.4%</td>
<td>93.1%</td>
<td>0.001</td>
</tr>
<tr>
<td>5-Year Subsequent Freedom from CAV</td>
<td>81.0%</td>
<td>89.7%</td>
<td>0.189</td>
</tr>
<tr>
<td>5-Year Subsequent Freedom from NF-MACE</td>
<td>66.7%</td>
<td>89.7%</td>
<td>0.039</td>
</tr>
<tr>
<td>1-Year Subsequent Freedom from ATR</td>
<td>66.7%</td>
<td>72.4%</td>
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</tr>
<tr>
<td>1-Year Subsequent Freedom from ACR</td>
<td>90.5%</td>
<td>86.2%</td>
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<tr>
<td>1-Year Subsequent Freedom from AMR</td>
<td>76.2%</td>
<td>82.8%</td>
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</table>

### Abstract 86

**Correlation of Complement Binding Donor Specific Antibody and Subsequent Outcome**

A Shen*, K Nishihara, R Levine, M Hamilton, J Kobashigawa. Smidt Heart Institute, Los Angeles, CA

10.1136/jim-2019-WMRC.85

**Purpose of study** It has been reported that the development of donor specific antibody (DSA) is correlated to the subsequent development of cardiac allograft vasculopathy (CAV, as defined by stenosis ≥30% by angiography) following heart transplantation. We now have the ability to assess whether DSA can bind complement (C1q+ assay) which suggests that these DSA are potentially cytotoxic to the donor heart and can result in CAV. We sought to assess if C1q+ DSA leads to the development of CAV in heart transplant patients.

**Methods used** Between 2010 and 2014, we assessed 50 heart transplant patients who developed DSA after heart transplantation. Patients were divided into groups that developed C1q+ DSA (n=21) and those that did not have this ability (n=29).

**Summary of results** Patients with C1q+ DSA had decreased freedom from non-fatal major adverse cardiac events (NF-MACE: myocardial infarction, percutaneous coronary intervention/angioplasty, new congestive heart failure, pacemaker/implantable cardioverter-defibrillator placement, and stroke), and freedom from rejection (any treated rejection (ATR), acute cellular rejection (ACR), antibody-mediated rejection (AMR)).

**Conclusions** The development of C1q+ DSA appears to be correlated with poor clinical outcome. It is not clear if treating C1q+ DSA with desensitization therapy can improve outcomes.

### Abstract 86 Table 1

<table>
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<tr>
<th>Endpoints</th>
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</tbody>
</table>

**Cardiovascular Disease Risk Begins Early in Males with Klinefelter Syndrome**

N Morelli*, J Kaar, K Moreau, S Davis. University of Colorado Anschutz Medical Campus, Aurora, CO

10.1136/jim-2019-WMRC.86

**Purpose of study** Klinefelter Syndrome (XXY) has an incidence of 1 in 600 births each year. Men with XXY have higher morbidity and mortality from cardiovascular disease (CVD). Increased carotid intima-media thickness (cIMT) is associated with CVD risk and has been shown to be higher in men with XXY. cIMT has not been assessed in adolescents with XXY. The objective of this study was to investigate CVD risk in adolescents with XXY with the hypothesis that elevated cIMT would be observed in youth with XXY.

**Methods used** This was a cross-sectional pilot study in adolescents with karyotype 47,XXY. cIMT was measured using ultrasound in the supine position by standard protocol. A one-sample t-test was used to compare cIMT in our sample to published norms for adolescent males (0.45±0.04 mm). A two-sample t-test was used to compare cIMT between boys treated with testosterone to those untreated. Correlation was used to assess relations between cIMT and clinical biomarkers of metabolic function, including seated systolic and diastolic blood pressure (SBP, DBP), pulse pressure, BMI percentile (BMIp), waist circumference (WC), fasting blood glucose (FBG) and lipid profile (total cholesterol, LDL, HDL, and triglycerides).

**Summary of results** cIMT in 25 males with XXY (mean age 14.5±1.5yrs, BMI percentile 55.3±32.6) was 0.55±0.07 mm, significantly greater than expected for this age (p<0.0001). There was no difference in cIMT in males on testosterone
Outcomes of Low-Birthweight Infants: Sustainable Improvement in Maternal-Neonatal Health Services Delivery in Rural Nepal

**Purpose of study** Low birthweight (LBW) babies (<2500 g) have a 20-fold higher mortality in their first year of life and account for 30% of neonatal deaths. In the Indian state of Gujarat, over 43% of infants are LBW. The purpose of this study is to determine the impact of a health surveillance intervention on growth and mortality in LBW infants in a rural area of Gujarat.

**Methods used** A pre-intervention assessment of LBW infants was conducted from January 6 to February 12, 2016. Infants were born at the Mota Fofalia Pediatric Center (MFPC) and discharged to villages in the Garudeshwar taluka. Families were not given newborn counseling or education. Assessment included weight and length.

A health surveillance intervention implemented in April 2016 provided pre-discharge counseling at the MFPC on infant warming and breastfeeding. Six follow-up household visits were made in the first year of life. Assessment included weight, length, temperature, and ECEB criteria for signs of infection. Families received counseling on infant care at each assessment.

**Summary of results** On baseline assessment, 72 of 86 identified infants were found for follow up (53% female; mean [range] age at follow-up, 4.7 [1–12] months). In the intervention phase, 330 LBW infants' families received pre-discharge counseling and 254 (77%) completed follow-up (56% female). There was no significant difference in weight gain between baseline (19.5 g/day) and intervention groups (19.0 g/day, t (360)=0.49, p=0.62). One-year infant mortality was lower in the intervention group (n=14, 4%) compared to baseline (n=8, 11%), as was neonatal mortality (intervention, n=3, 1%, baseline, n=3, 4%), though neither were significant (intervention, p=0.096, baseline, p=0.097).

**Conclusions** A health surveillance intervention in Garudeshwar with counseling on best feeding and care practices decreased LBW infant mortality. Future iterations of this study should address methods for increasing participant retention.

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**Conclusions**

This sample of normal-weight adolescents with XXY had a higher cIMT than would be expected for healthy males their age, suggesting their risk for CVD begins early despite the absence of obesity, hypertension, or diabetes. cIMT did not consistently correlate with common markers of cardio-metabolic health; however, this pilot study was underpowered for these analyses. Future studies with larger sample sizes, controls, and longitudinal assessments will help determine the pathogenesis and ultimately lead to earlier prevention and intervention for CVD in XXY.

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**Purpose of study** The purpose of this study is to describe the features and characteristics of an effective and sustainable multifaceted maternal-neonatal health intervention in rural Nepal.

**Methods used** Between 2015–2019, a public-private-academic partnership team implemented a multifaceted MCH intervention to build care capacity in the Humla and Solukhumbu districts of Nepal. Fourteen birthing center facilities were established at health posts, engaging the local government health system, local administrators, and the ministry of health to develop a sustainable model for improved MCH services delivery in existing rural clinics. We describe administrative, educational, and management interventions which led to self-sustaining improvements in service delivery for women and children. Fifteen skilled birthing attendants were provided with training, as well as 95 community-based providers for community liaison work and 75 health facility staff for basic perinatal care. Measures of care quality were measured using a standardized WHO survey before and after intervention.

**Summary of results** Deliveries at health facilities improved from 37% (before intervention) to 49% (after), deliveries attended by trained professionals increased from 27% to 46%. Women receiving postpartum hemorrhage chemoprophylaxis increased from 31% to 43%, delivery planning increased from 76% to 95%, and newborns receiving most or all essential elements of newborn care increased from 32% to 58%.

**Conclusions** Key features of the public-private academic partnership intervention included: 1) Local ownership, fingerprinting and accountability, local input into design, and local financial contributions to staff salaries and performance monitoring; 2) Improvement of case management capacity of staff through problem-focused hands on training experiences and government certification programs; 3) One-time investment in necessary equipment upgrades; 4) Integration of birthing centers within the government health system to access governmental supply chain for consumable materials; 5) Building local capacity for monitoring service delivery, care quality and outcomes through data systems and feedback loops.
EDUCATING DHULIKHEL HOSPITAL EMERGENCY DEPARTMENT PATIENTS AND FAMILY MEMBERS ABOUT THE RISK FACTORS OF CARDIOVASCULAR DISEASE

S Spooner*, University of Washington School of Medicine, Seattle, WA

Purpose of study The World Health Organization estimates that deaths due to noncommunicable diseases (NCDs) in Nepal have risen from 51% in 2010 to 60% in 2014, with 22% attributed to cardiovascular disease (CVD). Within the Nepali population there is limited education about CVD and its risk factors. In the absence of primary care, the first contact for patients and family members is often the Emergency Department (ED). A 2013 WHO cross-sectional study identified insufficient fruit and vegetable intake, obesity and overweight, hypertension, and hyperlipidemia as the main risk factors within the Nepali population.

Methods used A partnering Dhulikhel Hospital ED physician identified the need for CVD educational materials in the ED. A literature review of CVD in Nepal was conducted to identify the main risk factors locally. With the assistance of ED physicians and a research assistant, a poster was created featuring Nepali individuals and culturally relevant displays of risk factors. After feedback from ED staff and patients, the poster was revised to include common symptoms of CVD along with the Dhulikhel Emergency Medical Services (DEMS) contact information. The posters will be mounted and displayed on the walls in the ED and waiting room.

Summary of results Before printing, feedback from 10 ED patients or family members confirmed that the translation and grammar were correct, and the pictures depicting risk factors were clear. After printing, the poster was evaluated by the 13 ED staff for effectiveness using a short survey. On a scale of 1–5, with 5 being easy and 1 being very difficult to understand, 7 staff rated the poster at 4/5 with the remaining 6 rating it as 5/5. All 13 staff members thought a patient without a health background would be able to comprehend the poster’s meaning and stated that they would use it to educate their patients about CVD.

Conclusions The project succeeded in creating a culturally relevant educational platform for ED patients and their family members to learn more about the risk factors for CVD in Nepal. It also provides a description of symptoms associated with CVD, a visual of the DEMS contact information, and a reminder encouraging patients to seek care immediately. More work is needed to create materials educating patients about CVD risk-reducing behaviors.

ENGAGING TRADITIONAL MEXICAN MIDWIVES IN THE USE OF BAG VALVE MASKS TO TREAT ASPHYXIATED NEONATES

PF Draper*, University of Utah School of Medicine, Salt Lake City, UT

Purpose of study This project launched a collaborative partnership between an American medical school and a traditional midwifery school in Oaxaca, laying the groundwork for future neonatal health endeavors in partnership with traditional midwives (‘Parteras’) in the region.

Methods used In March 2019, two certified Helping Babies Breath master trainers—a neonatologist and a medical student—taught an 8-hour course for eighteen Parteras involving didactic content and several hours of hands-on practice with neonatal mannequins and bag valve masks.

Summary of results Participating Parteras clarified their ability to identify neonatal asphyxiation, gained insight into the strengths and weaknesses of their current interventions and learned to effectively utilize a bag valve mask to treat asphyxiated neonates.

Conclusions Parteras from across Oaxaca gained a tangible skill to address neonatal asphyxiation in low-resource settings. Several neonatal mannequins and bag valve masks were donated to the school to ensure ongoing training on this topic. As training is sustained by the midwifery school, traditional midwives will be equipped with the skills, resources, and confidence to identify areas of local need and implement their own training programs for other traditional birth attendants. This Training Of The Trainer model opens up long-term potential for reduced neonatal mortality due to asphyxiation in the rural communities of Oaxaca.
Purpose of study Establish baseline data for a rural hospital in Gujarat, India, quantifying intake vital sign assessment and qualitative information regarding care in the Emergency Department.

Methods used This direct observational assessment took place at Shree Chhotubhai A. Patel Hospital, a rural hospital in Gujarat, India. In January – July of 2019, medical students trained in data collection directly observed Emergency Care delivery and triage assessment using a standardized evaluation tool. The data collected is descriptive.

Summary of results A total of 43 patient encounters were assessed during the data collection period. Heart rate and peripheral capillary oxygen saturation (SpO2) were the most frequently assessed vital signs, measured in 30% (13/43) and 27% (12/43) of patient encounters, respectively. Blood pressure was measured for 25% (11/43) of patients. Least frequently measured was patient temperature at 7% (3/43) and respiratory rate at 2% (1/43).

The presenting symptoms for 47% (20/43) patients seen were for trauma, 12% (5/43) for obstetrics care, 7% (3/43) with concern for stroke, 5% (2/43) with chest pain. In terms of Emergency room staff availability, 70% patients were seen by nursing within 5 minutes of arrival to the emergency department, with maximum wait time of 20 minutes until a nurse was able to assess the patient. 70% (30/43) patients were seen by the medical officer on duty within 30 minutes of arrival.

Conclusions WHO guidelines recommend triage systems based on presenting signs and vitals. Staff are available to assess patients rapidly upon arrival to the emergency department. There is a need to improve the measurement of vital signs. One key element contributing to the challenge of a triage assessment is the limited availability of equipment needed for accurate assessment. Availability of pulse oximeters and mobile blood pressure cuffs led to a relative increase of vital sign measurement. There was not a working thermometer in the hospital. This baseline analysis helps initiate an effort to understand the current practices in the ED and informs the creation of plans to improve appropriate patient care delivery.

Improving maternal and child health are central to the realization of National Health Goals outlined in the National Rural Health Mission (NRHM). Key interventions include increased support of deliveries by skilled birth attendants (SBAs). This study provides a descriptive assessment of current maternal care delivery by SBAs in the peripartum period.

Methods used This assessment took place at Shree Chhotubhai A. Patel Hospital in rural Gujarat, India. In July, 2019, medical students trained in data collection directly observed perinatal care delivery to estimate SBA proficiency. SBA competency was assessed by medical students via a standardized evaluation tool, the components of which were adapted from the World Health Organization’s (WHO) top priority quality indicators.

Summary of results A total of 26 partial or complete care encounters were assessed during this data collection period. Of the 9 observed admissions, maternal blood pressure, temperature, and heart rate were assessed 44%, 0%, and 33% of the time, respectively. In these women, fetal heart rate was assessed 100% of the time, but was never documented. A total of 9 cesarean sections (C/S) and 16 vaginal deliveries were observed. Of the 16 vaginal deliveries, 11 (69%) women experienced inappropriate fundal pressure applied by SBAs. Of the 25 deliveries (16 vaginal, 9 C/S), 19 women (76%) received oxytocin at some point after delivery; only 6 women (24%) appropriately received oxytocin within 1 minute of birth.

Conclusions This evaluation found variation and deficiencies in perinatal care services provided by SBAs. There is a need to refine provider training and protocols to improve the quality of perinatal care.

Purpose of study The WHO ranks depression and anxiety as the two largest contributors to disability globally, with over 80% of affected people living in low- and middle-income countries. However, these conditions often remain undiagnosed and undertreated in developing countries due to stigma, sociocultural variance in perceptions of mental health, and limited regionally-specific knowledge. A greater effort to describe and study mental illness beyond a Western context is much-needed. The University of British Columbia’s Global Health Initiative (GHI) has a longstanding partnership with Munsell’ing Boarding School in northern India. After hearing anecdotal reports of high stress levels among students, the GHI team developed a preliminary study to screen for...
AN ANTHROPOMETRIC ASSESSMENT OF CHILDREN UNDER 5 IN RURAL KENYA

J B Creelman*, J Jacob, SM Gunn, C Pang, P Mbullo, V Kapoor. 1University of British Columbia, Vancouver, BC, Canada; 2Northwestern University, Evanston, IL

Purpose of study Globally, 1 in 9 people face hunger. 52 million children are wasted and 155 million have stunted growth. Pamoja, a community-based organization in the Kisumu district of rural Kenya, has programs to improve health, income and food security of its residents. In order to identify how best to design and deliver programs, Pamoja identified 434 children under age 5. Height-for-age (HAZ), weight-for-age (WAZ), and height-for-weight (HWZ) were measured to gauge stunting, underweight, and wasting, respectively. With written informed consent from the household head, community health workers collected anthropometric data. Supervision and the WHO training course on Child Growth Assessment were provided. Ethical approval was granted by UBC BREB and AMREF. WHO Anthro Analyzer software was utilized to calculate age-adjusted Z-scores, comparing to WHO Child Growth Reference data. Stunted, underweight, and wasted children were identified by Z-score values of ≤–2SD (moderate) and ≤–3SD (severe) based on HAZ, WAZ, and HWZ, respectively.

Summary of results 62 (14.3%) children had moderately stunted growth and 40 (9.2%) had severely stunted growth. 24 (5.5%) children were moderately underweight and 24 (5.5%) were severely underweight. 20 (4.6%) children had moderately wasted growth and 13 (2.9%) had severely wasted growth. 7 children (1.6%) had MUAC measurements in the moderate to severe malnourishment range and were referred to acute medical care.

Conclusions The most prevalent form of malnutrition in this study population is stunting, which is typically indicative of chronic nutrient deficiency. This baseline data enables Pamoja to effectively target nutrient-deficient families and track their service delivery and impact. It is important to note that all children included in this study are beneficiaries of Pamoja. Therefore, this study population was not randomly sampled, which presents selection bias and a limitation to interpretation.
in the allocation of funding and programs to improve the access to sufficient nutrition, and modify the food support to vulnerable families. The study is limited due to the reporting period of the HFIAS and HDDS. HFIAS captures data from the past month and HDDS from the past day. Thus, the average longitudinal consumption of a family might not be represented. Households will be followed with surveys and growth monitoring for 5 years.

96 THE UTILITY OF TELEDERMATOLOGY TO ENHANCE FOLLOW-UP CARE AND INFORMATION DISTRIBUTION IN LIMA, PERU


10.1136/jim-2019-WMRC.96

Purpose of study With the increasing use of WhatsApp among physicians and patients, this form of communication is being adapted to improve clinical practice. The purpose of this project was to identify ways to overcome barriers to care among patients in a dermatology clinic in Lima, Peru.

Methods used Fifty patient interviews were conducted to better understand healthcare barriers at the Hospital Nacional Daniel Alcides Carrion in Lima, Peru. The interviews also assessed WhatsApp use and if patients would be comfortable using it for follow-up care. In a pilot study, 10 patients were selected to receive follow-up care via WhatsApp. Over 8 weeks, patients sent in photos and concerns to a smartphone for this project. Responses were given by the chief dermatologist (EQ). To use WhatsApp for information distribution, educational materials were developed on three topics (self-medication, antibacterial soaps, and recommendations for skin products). Patients were asked about the readability and understanding of these messages. The 10 patients who completed WhatsApp follow-up care were asked about their comfort receiving dermatologic care via WhatsApp and whether they would do it again.

Summary of results From the patient interviews, common barriers included high cost, long travel times, and difficulty getting free time away from work or school. Seven (14%) of the 50 patients reported disabilities impacting travel. The majority (76%) of patients had smartphones capable of taking photos. From the pilot study, nine (90%) of the ten patients said they would do it again and all ranked their comfort as a 9 or 10 of 10. One patient did not give feedback. The response was overall positive, including comments like, ‘I want to thank you and Dr. Quijano for caring about my health. This means of communication is very useful.’

Conclusions In this pilot study, WhatsApp was a powerful tool that improved communication between physicians and patients, especially those who had difficulty attending in-person appointments. The hospital plans to continue using this method. In the future, we plan to compare outcomes in patients who receive follow-up via routine clinic visit or via WhatsApp. If outcomes are similar, WhatsApp could potentially provide a method to improve care for more patients.

Global health II
Concurrent session
3:15 PM
Thursday, January 23, 2020

97 IMPROVING UNDERSTANDING OF THE BURDEN OF TRAUMATIC INJURIES THROUGH TRAUMA REGISTRIES IN LIMA, PERU

T Jordan*, University of Washington School of Medicine, Seattle, WA

10.1136/jim-2019-WMRC.97

Purpose of study Trauma registries are databases that can be used to better understand the burden of trauma within a population. Data can be used to implement quality improvement projects, inform local policies, and develop targeted preventative health measures. Staff at Hospital de Vitarte in Lima have spent 13 months using a trauma registry, though much of their data is missing or incomplete. The goal of this project was to examine the process of data collection and entry and propose solutions to improve data quality.

Methods used Preliminary data were presented to hospital staff to illustrate current gaps. Meetings were held to gather feedback on how to improve data entry processes. Inclusion criteria were defined and will be posted on the wall of the department. A new paper intake form was created and a schedule was developed for interns to digitize intake forms. A training workshop was developed for and delivered to residents and interns. Training materials were left with staff for future workshops. Finally, a patient education pamphlet on pedestrian, bicycle, and automobile safety was developed for distribution after data analysis identified a high burden of traffic accident-related trauma.

Summary of results A total of 19 residents and interns were trained in the use of trauma registries and the new paper intake form. In a post-workshop survey, 90% of respondents indicated agreement that trauma registries are important tools for improving outcomes in trauma patients. Additionally, 84% of participants agreed that the new paper entry form would improve data quality and 79% thought that the new form would ease the process of online data entry. The Director of the Trauma Quality Committee of the Society of General Surgeons of Peru is seeking approval through that society and the Panamerican Trauma Society to distribute the patient education pamphlet as part of a larger public health campaign.

Conclusions This project led to the implementation of specific strategies for data quality improvement, including changes to the paper intake form, well-defined inclusion criteria, and weekly data upload and revisions. Additional funding and resources including dedicated administrative staff are needed to improve data entry and inform injury prevention strategies to reduce morbidity and mortality in trauma patients.
Purpose of study The pediatrics department of the Regional Hospital of Loreto has had 163 nosocomial infections reported in the last 5 years, and hand hygiene is an effective and inexpensive intervention to decrease interpersonal spread of microbes. This project aimed to improve hand hygiene through increased access to resources, reminders, and educational materials.

Methods used An initial assessment of the availability of hand hygiene resources and of staff and family hand hygiene compliance was performed. Together with quality control and pediatrics department staff, I worked to increase access to hand hygiene materials; place WHO reminders and education materials for hand hygiene in patient rooms, bathrooms, and sinks; and evaluate technique, compliance, and barriers to proper hand hygiene among staff and families to provide performance feedback. We provided a final written report of the department’s hand hygiene compliance status and recommendations for improvement to the office of quality control.

Summary of results The initial assessment identified a lack of hand hygiene resources throughout the department and low sanitation standards among personnel and patient families. Barriers to hand hygiene voiced by hospital staff included lack of resources, inconvenience from washing or dislike of hand sanitizer, and preferring to simply wear gloves. By the project’s end, all rooms were equipped with WHO hand hygiene educational materials and hand sanitizer, and bathrooms and sinks were equipped with soap and WHO educational materials. Most personnel demonstrated proper knowledge of technique, but few knew the WHO 5 moments when hand hygiene is indicated. Staff that were in the habit of cleaning their hands appreciated and utilized the increased access to hand hygiene materials and reported that they now more frequently clean their hands. Most families evaluated utilized the hygiene materials and reported that they now more frequently clean their hands. A t-test compared growth velocity and z-score changes from birth to discharge between infants with LOS ≤14 days vs. >14 days. Regression analysis predicted weight z-score change based on LOS. A p-value <0.05 was statistically significant.

Summary of results 283/704 (40.1%) infants were born ≤36 weeks GA. Mean birth GA was 36.6 weeks with weight 2.61 kilograms (kg) (z-score -0.024). Mean LOS was 7.4 days with discharge weight 2.54 kg (z-score -0.855). Infants with LOS >14 days (mean 24 days) experienced more growth failure than infants with LOS ≤14 days (mean 4 days) with change in z-score -1.66 vs. -0.64 (p<0.001). Growth velocity of infants with LOS >14 days was low at 4.7 grams/kg/day, with each day of hospitalization predicting a -0.044 decrease in weight z-score (p<0.001).

Conclusions Infants admitted to a newborn unit in a government hospital in Nakuru, Kenya experience growth rates that are inadequate to meet recommended growth velocities by the World Health Organization. Longer lengths of hospital stay contribute to decreases in weight z-score and an increased risk of malnutrition. Additional assessment and intervention is needed to promote improved nutrition provision and growth rates of these infants.

Purpose of study Inadequate nutrition remains a leading cause of growth failure in neonates, especially those born preterm or critically ill. Previously, members of this research team implemented a respiratory intervention in the newborn unit in a government hospital in Nakuru, Kenya. Infant growth failure in the unit was suspected based on clinical observation. The purpose of this study is to quantify growth velocity and weight z-score changes for infants in a Kenyan newborn unit from birth to hospital discharge.

Methods used After ethical approval, data was collected from the hospital charts of Kenyan infants (n=704) admitted to a newborn unit between June 2016 to December 2018. Collected information included birth gestational age (GA), birth weight, discharge weight, length of hospital stay (LOS) in days, and z-scores for weight on the INTERGROWTH 21st growth chart. Infant growth velocity was calculated as follows:1,000/ln( discharge weight/birth weight)/LOS. Averages are reported in means. A t-test compared growth velocity and z-score changes from birth to discharge between infants with LOS ≤14 days vs. >14 days. Regression analysis predicted weight z-score change based on LOS. A p-value <0.05 was statistically significant.

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Purpose of study Pneumonia rates in Naivasha sub-county have doubled in the last 5 years, with a larger percentage of cases admitted to Naivasha Community District Hospital (NCDH) coming from the Karigita community. Current research reports the changing nature of pneumonia to predominantly viral causes in response to effective bacterial vaccination and lack of attention placed on primary prevention. Professional Community Health Education Workers (CHEW) proposed training Community Health Volunteers (CHV) to educate households during work for ongoing projects.

Methods used Building on ongoing local research, public health projects, and based on Kenyan and international guidelines, a training was developed to cover pneumonia-
related misconceptions and preventative methods. Ongoing public health interventions already address immunization, indoor pollution, and hygiene to prevent diarrheal diseases, so emphasis was placed on incorporating cough hygiene, addressing misconceptions about pneumonia, educating about prompt care-seeking, avoiding antibiotic misuse, and promoting exclusive breastfeeding for 6 months. CHVs participated in a 1-hour educational session with facilitated discussion and pre/post-knowledge-based assessments. A 2-month follow up and data tracking of pneumonia cases will be used to assess the quality of the training and its impact on the community.

Summary of results Thirty CHVs participated in this educational session, with the assistance of the Ministry of Health Public Health Branch, NCDH, and NIH funded research. The pre to post-test knowledge assessment score increased over 80% from 4/30 to 29/30. Continued education on pneumonia prevention is ongoing within Karijita and neighboring communities as a direct result of a successful educational session. A digital media for more efficient sharing, storage, and access was suggested and a readily accepted alternative to printed educational material.

Conclusions Participating CHVs benefited from an increase in knowledge-base and were actively engaged in discussions to address methods of sensitizing the community and implementing change in cultural practices. The integrative scope of the project and the session’s engaging atmosphere generated opportunities to collaborate with ongoing research and set-up additional workshops addressing community needs.

Summary of results At baseline (2014), healthcare providers did not appropriately identify neonates as low birth weight (LBW) (0%, n=26) and did not regularly provide appropriate inpatient monitoring (daily check of temperature, heart rate, respiratory rate) (0%, n=138).

Postintervention analysis (2019) data show that training gradually impacted obtaining and recording regular weight values. 67% of neonates (14/21) were appropriately identified as high-risk LBW infants at birth, and 48% of LBW patients were appropriately discharged (>1800 g, feeding well, 10/21); appropriateness of daily newborn assessment, including vital signs, improved to only 20% (9/44). Significant barriers to care quality include a lack of adequate personnel and the availability of functional equipment staff feels confident using.

Conclusions Despite educational and QI interventions, improvement in newborn care quality remains modest. Additional system strengthening interventions are needed.
the educational intervention. Further studies will be required to assess retention of knowledge and clinical translation to neonatal outcomes following educational interventions with HBB, ECSB curricula.

**Purpose of study**

Schistosome parasites cause a chronic inflammatory disease in humans. Past evidence suggests that these parasites can detect macromolecules of their snail vector hosts; however, it is unclear if they can differentiate among potential hosts. Understanding host seeking behavior could lead to novel control strategies to prevent human infection.

**Methods used**

Using wild strains of African snails and parasites, we tested the hypothesis that parasites are attracted to the most susceptible host. We tested attraction to three snail species: Biomphalaria sudanica (BS), B. choanomphala (BC), and B. pfeifferi (BP) using choice chamber experiments in which parasites could orient toward a given snail stimulus. Attractiveness was assessed visually via microscopy by counting parasites that moved into each chamber toward stimuli.

**Summary of results**

Analysis indicated that miracidia were significantly more attracted to snail-incubated chambers compared to empty control chambers, indicating an attraction to all three snail species. In pairwise comparisons, we found that in some cases, the parasites were significantly attracted to the more susceptible species (BC v. BS), but in others, there was no significant difference (BS v. BP). Finally, we found a significant aversion of parasites to snails already infected with a parasite that preys on schistosome larvae within the snail, suggesting that the schistosome parasites can actively detect and avoid these predators.

**Conclusions**

The results of these experiments indicate that schistosomes use chemical signals to find hosts, and in some cases can differentiate signals to orient toward a host that will maximize their chance of successful establishment.

**Purpose of study**

Following radiation for glioma, new enhancements on MRI present a challenge in differentiating recurrent tumor and radiation-induced lesions, termed pseudoprogression (Ps). Criteria have been outlined to diagnose Ps after photon radiation by the Radiologic Assessment in Neuro-Oncology (RANO) group based on time and location of occurrence. Some patients receiving protons manifest changes that seem subjectively different in appearance, location and timing from photon Ps, and would be identified as recurrence. We retrospectively review post-treatment MRI changes of proton patients and compare the Ps seen after photon radiation. We propose a criterion to characterize proton pseudoprogression (PRoPs) distinct from photon.

**Methods used**

Post-treatment imaging of patients with gliomas were reviewed, along with clinical and pathological data. 74 proton patients were reviewed for the presence of PRoPs, and then 64 photon patients were reviewed for any matching imaging changes. Data collected included the location, timing, and morphology of the imaging change, tumor grade, molecular subtyping and chemotherapy received.

**Summary of results**

Thirteen patients (16%) received protons and had imaging changes unique to treatment with protons, which we term PRoPs (Proton Pseudoprogression). We established the following criteria to characterize PRoPs:

1. Located not immediately in or adjacent to the resection cavity or residual tumor; located ~ 2 cm from target opposite proton beam entry; asymptomatic (unless simultaneous disease progression is also occurring); resolves without treatment, and some combination of subjectivity multifocal, patchy, small (<1 cm).

In the comparison group receiving photon, 0 had changes that met our criteria for PRoPs (p=0.001). The mean time when PRoPs occurred was 16.2 months after radiation.

**Conclusions**

Patients who receive protons are subject to a unique subtype of Ps that can mimic tumor progression. PRoPs can possibly be explained by the increased Relative Biological Effectiveness (RBE) of protons and beam angle selection. Current RANO guidelines would inaccurately characterize PRoPs as tumor progression. Using the radiation oncology treatment plan can help confirm the nature of the enhancement and prevent unnecessary treatment for mistaken tumor progression.
transcriptional reactivation of p16^{INK4A} can inhibit cell proliferation in cSCC.

**Methods used** We transduced cSCC cell line A431 with lentivirus containing nuclease-deactivated Cas9 (dCas9) fused to the catalytic domain of histone acetyltransferase p300 (dCas9-p300), which catalyzes acetylation of histone H3 at lysine 27 (H3K27ac). Alternatively, we transduced A431 with lentivirus containing dCas9 fused to transcriptional activator VP64 (dCas9-VP64). After drug selection to enrich transduced cells, we performed a second lentiviral transduction to introduce guide RNA (gRNA) directing dCas9 fusion proteins to the p16^{INK4A} promoter within the CDKN2A locus. After one week of further drug selection, we collected these cells and measured p16^{INK4A} mRNA expression (RT-qPCR) and cell proliferation (Cell Counting Kit-8).

**Summary of results** RT-qPCR revealed that dCas9-p300 and dCas9-VP64, each with gRNA targeting the p16^{INK4A} promoter within the CDKN2A locus, increased p16^{INK4A} mRNA expression in A431 cells ~30–40-fold relative to untransduced cells. Both dCas9-p300 and dCas9-VP64 targeting the p16^{INK4A} promoter reduced cell proliferation by ~20–30%. Increased p16^{INK4A} expression correlated with decreased cell proliferation in these modified cells.

**Conclusions** This study demonstrates that CDKN2A can be upregulated by targeted epigenetic modification or transcriptional activation, leading to inhibited cell proliferation. Future investigation will determine the kinetics, robustness, and off-target effects of targeted transcriptional reactivation tools. Targeted epigenome editing of cancer-relevant genes will be a versatile basis for precision anticancer therapies.

**DELCIVERY OF ADJUVANTS WITH C3-TARGETING LIPOSOMES LEADS TO MYELOID CELL ACTIVATION FOR CANCER IMMUNOTHERAPY**

D Heald*, M Kulberg, H Martinson. University of Washington, Anchorage, AK

**Purpose of study** Cancer immunotherapy treatments have gained much popularity recently, with many different pathways being targeted, including toll-like receptors (TLRs). These receptors have been implicated in tumor suppression via cytokine stimulation and as such, TLR agonists have shown promise in promoting tumor reduction in several cancer models. One such successful model is by use of liposomes to deliver these agonists. There is a lack of data, however, on what the exact mechanism is by which these TLR agonists are affecting tumors, which is important for understanding their safety and efficacy.

**Methods used** In healthy C57BL/6 mice, bone marrow cells were treated in vitro and then underwent qPCR to determine gene expression for specific cytokines. Treatments included PBS, naked control and C3 liposomes, and control and C3 liposomes with TLR agonists encapsulated. In BALB/c mice with breast cancer, an identical method was used for obtaining PCR data, but treatments differed. These treatments were PBS, control and C3 liposomes with free TLR agonists, and control and C3 liposomes with TLR agonists encapsulated. The TLR agonists utilized were MPLA, CpG, and R848, and the genes examined included IL-1β, IL-6, TGF-β1, IP-10, and IRF-7.

**Summary of results** In the C57BL/6 mouse, there was a significant increase in expression in the C3 with TLR agonist group for IL-1β, IL-6, IP-10, and IRF-7. Also, TGF-β1 expression was significantly reduced in TLR agonist encapsulated groups when compared to PBS. In the BALB/c mouse, the control and C3 with encapsulated agonist did show increased IL-6 expression, and these two groups along with the C3 and free agonist group showed increased IRF-7 expression. For IL-1β and IP-10, the four treatment groups did show increased expression compared to PBS.

**Conclusions** In the C57BL/6 mouse, delivering TLR agonists via a C3 liposome showed increased expression of anti-tumor cytokines, as well as decreased expression of TGF-β1, a pro-
Abstracts

**108** RADIOGENOMIC MODEL FOR DETECTING PROSTATE CANCER

N Oune*, MT Nguyentat, A Sasani, P Bhattar, C Chahine, PA Kohanteb, B Blome, E Uchio, R Houshyar. University of California Irvine Medical Center, Orange, CA; University of Colorado Anschutz Medical Campus, Aurora, CO

Purpose of study Multiparametric MRI (mpMRI) utilizing Prostate Imaging Reporting and Data System version 2 (PI-RADSv2) can accurately detect prostate cancer (PCA). Genomic data shows overexpression of Prostate Cancer Gene 3 (PCA3) in patients with PCA. However, no studies integrate these 2 indices. We propose a radiogenomic model amalgamating PCA mpMRI and PCA3 to predict clinically significant PCA (csPCA).

Methods used 147 consecutive patients who underwent digital rectal exam, PCA3 testing, prostate mpMRI, and MRI/TRUS Fusion Biopsy were included. PI-RADSv2 criteria was utilized for mpMRI. csPCA was defined as Gleason Score ≥ 7. Logistic regression, odds ratio (OR), and Area Under the Curve (AUC) were performed for correlation and accuracy.

Summary of results For a total of 266 lesions we included 3 independent predictors of csPCA: PIRADSv2 (OR 3.9, 95% CI 2.5–6.0), DRE (OR 2.6, 95% CI 1.2–5.5), and PCA3 (OR 1.0, 95% CI 1.0–1.0). The specificity, negative predictive value (PV), sensitivity, and positive PV achieved were 91%, 82%, 50%, and 31%, respectively. Our model’s AUC (0.81, 95% CI 0.75–0.87) was greater than that of PIRADSv2 (0.77, 95% CI 0.71–0.84), PCA3 (0.69, 95% CI 0.61–0.76), and DRE (0.56, 95% CI 0.48–0.64) alone.

Abstract 108 Figure 1 ROC curve

Conclusions A combined PCA3 and mpMRI radiogenomic model demonstrates improved AUC for predicting csPCA. Applying this model in clinical practice may decrease unnecessary biopsies and reduce cost and morbidity.

**109** ACTIVATION OF PROTEASOME BY INHIBITING AUTOPHAGY IN CORNEAL EPITHELIA CELLS WITH LIMBAL STEM CELL DEFICIENCY

D Carter*, F Bardag-Gorce, N Nilihara. The Lundquist Institute Harbor-UCLA, Torrance, CA; Charles R. Drew University of Medicine and Science, Los Angeles, CA

Purpose of study Previously, failure of UPP (Ubiquitin Proteasome Pathway) was found associated with an activation of autophagy in corneal epithelial cells with limbal stem cell deficiency (LSCD). However, autophagy activation does not remove damaged, aggregated proteins such as keratins (K4 and K13). The present hypothesis proposes that UPP activation might increase the clearance of misfolded proteins including K4 and K13, improving corneal epithelial cell function.

Methods used Rabbits with surgically-induced LSCD were used to quantify the elements of both pathways in corneal epithelial cells (CEC). Rabbit oral mucosa epithelial cells (OMECs) as, similar to conjunctival epithelial cells, are rich in K4 and K13. OMECs were isolated, cultured and treated with proteasome inhibitors and chloroquine to inhibit autophagy. Morphologic analysis of corneal tissue sections showed that both pathways stained positive in normal corneal epithelium.

Summary of results While constitutive proteasome beta subunits B1, B2 and B3 were decreased, autophagy biomarkers – ATG5 and MAPLC3B, were significantly increased in LSCD-CEC. However, despite autophagy up regulation, modified K4 and K13 still deposited and accumulated in LSCD-CEC without clearance. Proteasome inhibition in OMECS also showed a significant increase in ATG12, ATG5 and MAPLC3, confirming our observation that when UPP is defective, autophagy is stimulated. Additionally, when autophagy was inhibited in OMECS using chloroquine, our results showed not only an increase in proteasome chymotrypsin-like activity, but also a significant decrease in unmodified K4 and K13 levels with no keratin high molecular weight deposition.

Conclusions Proteasome activation in the ocular surface could be used to alleviate corneal epithelial cell dysfunction associated with LSCD.

**110** DEEP LEARNING FOR PROSTATE SEGMENTATION: CASE VOLUME AND PERFORMANCE

J Carbone*, M Bardis, A Sasani, C Chahine, P Bhattar, H Liu, P Chang, R Houshyar. University of California Irvine Medical Center, Orange, CA

Purpose of study Deep learning analysis of medical images is a data hungry challenge. However, few studies have systematically calculated the ideal number needed to balance training and performance. Here, we examine this question through quantifying the neural network (NN) performance of prostate organ segmentation against the number of samples.

Methods used This IRB approved retrospective review includes patients who had a prostate mpMRI between 9/2014 and 8/2018 and a MR-guided transrectal biopsy. A board-certified abdominal radiologist manually segmented each prostate organ on the T2 weighted images (T2WI). This study used our
customized Hybrid 3D/2D U-Net architecture and kept the same hyperparameters across the different NN runs. A total of 400 patients were available for training and validation. This study only changed the number of accessions fed into the NN and overall dice score was calculated with these patient numbers: 8, 16, 24, 32, 40, 80, 120, 160, 200, 240, 280, 320. During validation, the maximum number of available patients was used for the overall dice score calculation i.e. when the NN trained in 40 patients, the validation was completed on 360 patients.

Summary of results This study used 400 patients’ T2WI. The overall dice score varied from 0.424 with 8 patients to 0.867 with 320 patients with 0.800 reached at 80 patients (figure 1).

Conclusions The Hybrid 3D/2D U-Net for prostate segmentation produced the highest overall dice score of 0.867 at 320 patients. The performance improved most significantly when the patient number changed from 8 to 16, and it started to plateau at 160 patients. The performance only improved by 0.09 from 160 to 320 patients.

Abstract 110 Figure 1 Overall dice score vs. number of accessions
The overall dice score had its best performance of 0.867 at 320 patients. The biggest increase in performance occurred when increasing patient size from 10 patients to 20 patients. The performance started to plateau at 160 patients.

Summary of results In assessing the top 100 websites on glioma, it was found that 53% of the websites were commercial, and 7% of the websites were affiliated with an academic institute. Less than half of the websites provided authorship (40%) and cited sources (48%). About half (52%) stated the date of website creation, but only 38% of the websites identified dates of modification to content. A total of 26% of the websites had been updated within the last two years, with 66% of the websites last updated over four years ago. The majority of websites provided users with an in-site search engine as an interactive tool (86%), however, only 12% provided a discussion board or forum. In assessing readability, only 5% of the websites were readable at a level below grade 8 (elementary level). Almost all websites covered definition (91%) and treatment (92%), but very few of the websites covered prevention (4%). A total of 4% of the websites covering definition, and 4% covering treatment, provided inaccurate information.

Conclusions Analysis of the top 100 websites relating to glioma revealed there is a lack of current and understandable information available to patients online. A lot of the information available is not reliable due to a lack of cited author(s) and sources. Although the majority of sites covered definition and treatment of glioma, many other topics were poorly covered, including etiology/risk factors, prevention, and prognosis.

111 VETTING THE QUALITY OF AVAILABLE ONLINE INFORMATION FOR PATIENTS WITH GLIOMA
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10.1136/jim-2019-WMRC.111

Purpose of study The internet has become an invaluable source of information, both easily accessible and widely used. However, the quality of information online cannot be guaranteed. Glioma is the most common of all CNS neoplasms, yet little work has been done to explore the extent and quality of information online. Physicians must be aware of available online information to identify errors and gaps to better guide their patients’ self-directed glioma research.

Methods used The term ‘glioma’ was searched in Google and the meta-search engines Dogpile and Yippy, resulting in over 12.5 million hits. The top 100 websites averaged from these three search engines relating to patient information were analyzed using a structured rating tool. The websites were assessed based on affiliations, accountability, interactivity, site organization, readability, and content quality.

Abstracts
CUTANEOUS METASTASIS OF UROTHELIAL CARCINOMA RESULTING IN VASCULAR OCCLUSION AND LIVEDO RACEMOSA

1A. Swell*, 1B. Morris, 3M. Heaphy, 1University of Nevada, Reno School of Medicine, Reno, NV; 3Skin Cancer and Dermatology Institute, Reno, NV
10.1136/jim-2019-WMRC.112

Introduction A skin metastasis can be the first sign of an underlying visceral neoplasm and clinical presentation can be highly variable, making clinical diagnosis of these metastases challenging. Follow up investigations are often required to establish the location of the primary. Cutaneous metastases are a known but uncommon manifestation of urothelial carcinoma.

Livedo reticularis is a physical sign of the skin which presents as a net-like pattern consisting of macular, violaceous rings. Livedo racemosa is a pathologic form of livedo reticularis which is often associated with vaso-occlusive disorders. Both livedo reticularis and livedo racemosa arise secondary to dilation of the skin’s venous plexus which may be due to vasospasm, vasculopathy, or coagulopathy. Few cases of livedo reticularis or racemosa related to malignancy have been reported in literature and none describe vascular obstruction by tumor thrombus as the source of livedoid pattern.

Case report In this report, we describe an 82-year-old male with a history of urothelial carcinoma who presented to clinic with multiple, erythematous nodules in his right groin and a livedo-reticularis-like rash covering the anterior surface of his right thigh. Biopsies were taken from nodules in the right groin and livedoid patches on the right thigh. Those taken from nodules in the right groin were positive for cytokeratin 7 (CK7) and cytokeratin 20 (CK20), consistent with the original pathology report that diagnosed urothelial carcinoma and metastatic deposits in a right-groin lymph node. Biopsies taken
from livedoid patches on the right thigh were also CK7+/ CK20+; but more remarkably, microscopic investigation of the tissue and staining with endothelial cell markers CD31 and CD34 showed intravascular tumor thrombus as the source of vascular occlusion causing the livedoid pattern.

**Implications** This case report discusses the rare occurrence of cutaneous metastases of urethelial carcinoma and the novel finding of intravascular occlusion by metastatic carcinoma as the cause of livedo racemosa.

**Summary of results**

**Case 1:** Patient is a 63-year-old female diagnosed with Stage IVA serous carcinoma of the ovary with a history significant for prior breast augmentation. Initial diagnosis was made after presenting to the emergency department complaining of shortness of breath. A CT of the chest demonstrated a 16 × 14 × 16 cm pelvic mass and right sided pleural effusion. Thoracentesis identified malignant ovarian cancer cells. The patient underwent surgical debulking and completed eight cycles of chemotherapy. On physical exam after surgical debridement, it was noted the scar on the right breast from the breast implant procedure was distorted and hard. Further evaluation showed metastatic ovarian cancer in the surgical scar. After chemotherapy completion, she was started on maintenance therapy with Avastin due to increased tracer uptake was noted.

**Case 2:** Patient is a 56-year-old male initially diagnosed with prostate adenocarcinoma Stage II and history significant for prior chest pain. Initial treatment included 15 cycles of radiation along with Lupron Depot and Casodex for 3 months. Due to the side effects on the Lupron, the patient discontinued the Lupron with close follow up of PSA levels to monitor for recurrence. Nine months after stopping the Lupron, his PSA levels began to increase and the patient began to experience worsening back pain. An MRI of the spine was obtained due to the back pain which demonstrated potential prostate cancer recurrence with metastatic lesions noted at the L4 vertebrae.

**Conclusions** Locus Minoris Resistentiae is an important to understand in the setting of metastatic cancer. Identifying areas of previous tissue compromise might lead to earlier identification of metastatic disease. This could potentially lead to earlier and more effective treatment and management of such cases.

**Purpose of study** Locus Minoris Resistentiae is a phenomenon in which previously damaged tissue is less resistant to microorganisms. This is noted to occur in infectious processes but has not been well documented in the setting of cancer metastasis. These cases demonstrate the potential of metastatic preference in cancer to previous areas of tissue compromise with less resistance than other healthier locations in the body.

**Methods used** Retrospective case reports.

**Summary of results**

**Case 1:** Patient is a 63-year-old female diagnosed with Stage IVA serous carcinoma of the ovary with a history significant for prior breast augmentation. Initial diagnosis was made after presenting to the emergency department complaining of shortness of breath. A CT of the chest demonstrated a 16 × 14 × 16 cm pelvic mass and right sided pleural effusion. Thoracentesis identified malignant ovarian cancer cells. The patient underwent surgical debulking and completed eight cycles of chemotherapy. On physical exam after surgical debridement, it was noted the scar on the right breast from the breast implant procedure was distorted and hard. Further evaluation showed metastatic ovarian cancer in the surgical scar. After chemotherapy completion, she was started on maintenance therapy with Avastin due to increased tracer uptake was noted.

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**Conclusions** Locus Minoris Resistentiae is an important to understand in the setting of metastatic cancer. Identifying areas of previous tissue compromise might lead to earlier identification of metastatic disease. This could potentially lead to earlier and more effective treatment and management of such cases.
115 UTILIZATION OF FUNCTIONAL ECHOCARDIOGRAPHY AND NEAR-INFRARED SPECTROSCOPY TO CHARACTERIZE DUCTAL HEMODYNAMICS IN THE PREMATURE NEONATE

M Ringle*, VY Chock, K Stauffer, L Lopez, R Punn, S Bhombal. Stanford University, Palo Alto, CA

Purpose of study Echo is used for determining a hemodynamically significant duct (hsPDA) in the preterm neonate. Near-infrared-spectroscopy (NIRS), a noninvasive modality provides continuous feedback of organ perfusion. Historically, echo is used to diagnose ductal patency, and NIRS used for perfusion monitoring; rarely have they been used together to assess the duct. We aim to use the combined technology of echo and NIRS to identify patients with a hsPDA, as well as explore the possibility of solely using NIRS to identify and continuously monitor patients with a hsPDA.

Methods used Infants born <30 weeks at the Packard Children’s Hospital NICU are enrolled for echoes on DOL 3, 5, 7, 14, and 30. Cerebral and renal NIRS are obtained with each echo. A hsPDA is based on measurements of ductal characteristics, markers of pulmonary overcirculation, and systemic perfusion.

Summary of results We have 12 patients and 51 echoes with NIRS correlates; 18% show a hsPDA. The mean renal NIRS for infants with a hsPDA was 54 compared with 68 (p=0.001) for infants without a hsPDA. Cerebral NIRS for infants with a hsPDA was 60 vs 72 (p=0.0005) without. All infants with hsPDAs were born <27 weeks gestation and with a birth weight <1000 grams. We did not find any significant difference in mode of respiratory support or enteral feeding patterns.

Conclusions When correlated with echo, we conclude that NIRS may be an adequate surrogate for a hsPDA, with renal NIRS showing greater variability, and should be utilized in neonates who are at high risk for a hsPDA.

116 RANDOMIZED TRIAL COMPARING EFFECT OF DIFFERENT EPINEPHRINE DOSES AND FLUSH VOLUMES IN A PERINATAL MODEL OF ASPHYXIAL ARREST

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Purpose of study Current NRP guidelines recommend administration of 0.5–1 ml flush following epinephrine dose (0.01–0.03 mg/kg) via low umbilical venous catheter (UVC). This flush volume may not be adequate for epinephrine to reach right atrium and there are no pharmacokinetic/efficacy/safety studies comparing 0.01 vs. 0.03 mg/kg of epinephrine via low UVC.

Methods used Asystole was induced by umbilical cord occlusion and resuscitation was initiated after 5 min of asystole. After 5 min of resuscitation per NRP guidelines, lambs were randomized to receive either 0.01 mg/kg (low dose) or 0.03 mg/kg (high dose) of epinephrine via low UVC.
REVISION OF TRACHEAL SUCTION GUIDELINES FOR INFANTS BORN THROUGH MECONIUM STAINED AMNIOTIC FLUID (MSAF): CHANGING CLINICAL SIGNIFICANCE OF ‘NON-VIGOROUS’ STATE

Purpose of study In 2016, the Neonatal Resuscitation Program (NRP) recommended that resuscitation of non-vigorous infants with MSAF follow the same principles as for those with clear fluid. Studies have shown increased respiratory morbidity among these infants after implementation of new guidelines.

We hypothesize that babies who remain non-vigorous despite stimulation represent a sicker cohort of infants compared to non-vigorous babies immediately after birth undergoing routine tracheal suctioning.

Methods used Detailed review of the hospital course of all infants born with MSAF at UC Davis Medical Center between Dec 2013- Feb 2014 (Era-1) was compared to infants born between Jan-April 2014 (Era-2).

Summary of results Records of 428 and 461 consecutively born neonates before and after implementation of new NRP guidelines were screened. Incidence of MSAF (10.3% vs 10.2%) and characteristics were similar (table 1). In era-1, 8 babies were labeled non-vigorous and all underwent tracheal suction. Only one infant (29 week gestation) required admission and respiratory support. In era-2, 4 babies were non-vigorous despite stimulation and all required NICU admission and respiratory support.

Conclusions ‘Non-vigorous’ infants born through MSAF in the routine-suction era were less sick compared to persistently non-vigorous infants despite stimulation in era-2. Future studies should use MSAF as the denominator (and not non-vigorous) while comparing the outcomes.

Abstract 117 Table 1 Values presented as n (% of MSAF neonates) OR median (IQR)

<table>
<thead>
<tr>
<th></th>
<th>Era 1</th>
<th>Era 2</th>
<th>p value</th>
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<tbody>
<tr>
<td>Total number of neonates born during the period</td>
<td>428</td>
<td>461</td>
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<tr>
<td>Number of neonates with MSAF</td>
<td>44 (10.3%)</td>
<td>47 (10.2%)</td>
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<td>Cesarean section</td>
<td>9 (20.5%)</td>
<td>9 (19.1%)</td>
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<td>Gestational age (weeks)</td>
<td>40 (39-40)</td>
<td>39 (38-40)</td>
<td>0.6</td>
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<td>Birth weight (grams)</td>
<td>3445 (3180-3670)</td>
<td>3420 (3120-3660)</td>
<td>0.007</td>
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<tr>
<td>Male</td>
<td>24 (54.5%)</td>
<td>22 (46.8%)</td>
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<tr>
<td>1 minute Apgar ≤3</td>
<td>8 (18.2%)</td>
<td>0</td>
<td>0.007</td>
</tr>
<tr>
<td>Endotracheal intubation for suctioning</td>
<td>8 (18.2%)</td>
<td>1 (2.1%)</td>
<td>0.027</td>
</tr>
<tr>
<td>Non vigorous</td>
<td>8 (18.2%)</td>
<td>4 (8.5%)</td>
<td>0.3</td>
</tr>
<tr>
<td>Respiratory distress requiring NICU admission</td>
<td>2 (4.5%)</td>
<td>8 (17.7%)</td>
<td>0.11</td>
</tr>
<tr>
<td>Total admission to NICU/CICU</td>
<td>4 (9.1%)</td>
<td>11 (23.4%)</td>
<td>0.12</td>
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<tr>
<td>Non vigorous babies (%)</td>
<td>8</td>
<td>4</td>
<td>0.02</td>
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<tr>
<td>Required admission*</td>
<td>1 (12.5%)</td>
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<td>Highest Resp support after admission</td>
<td>12.5%</td>
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<td>- Mechanical Ventilation*</td>
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<tr>
<td>- HFNC*</td>
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</table>

*Percentage of non vigorous neonates
FACTORS ASSOCIATED WITH ADVERSE EVENTS AND VARIATIONS IN CAFFEINE USE FOR APNEA OF PREMATURITY IN PRETERM INFANTS IN THE UNITED STATES

1E Sander*, 1I Shepherd. 1LAC+USC Medical Center, LA, CA; 2CHLA, LA, CA

10.1136/jim-2019-WMRC.120

Purpose of study The Caffeine for Apnea of Prematurity trial demonstrated that caffeine administration within the first ten days of life in extremely premature infants to treat apnea led to decreased rates of bronchopulmonary dysplasia, death, and neurodevelopmental disability. Recent retrospective studies have suggested that caffeine used prophylactically can lead to similar improved outcomes; however, there are no known large prospective studies proving the effectiveness and safety of prophylactic caffeine and no consensus that prophylactic caffeine should be used. We surveyed NICU medical directors in the United States to delineate variations in usage of caffeine in order to identify potential focuses for future research and guidelines development.

Methods used An online survey with questions regarding various aspects of caffeine use was emailed to all NICU medical directors in the United States.

Summary of results 127/532 (24%) of medical directors completed the survey. The majority are at level III or IV NICU’s (95%) and are academically affiliated (67%). The most common reasons for initiation of caffeine include treatment of apnea (87%), prophylaxis for apnea (73%), and facilitation of extubation (59%). While 69% start caffeine soon after birth, only 34% discuss caffeine during the prenatal consult. Of those, only a third discuss potential side effects. Post menstrual age (PMA) (53%) is the most important factor in the decision to discontinue caffeine, followed by cessation of apneic episodes (42%). 34 weeks is the most common PMA to discontinue caffeine, followed by cessation of apneic episodes (42%). 34 weeks is the most common PMA to discontinue caffeine, followed by cessation of apneic episodes (42%). 34 weeks is the most common PMA to discontinue caffeine. Neonates are monitored for an average of 6 days following cessation of caffeine prior to discharge. Most (70%) do not send infants home on caffeine. Of those who do, 61% send those infants home with an apnea monitor.

Conclusions Caffeine is widely used in preterm neonates both early and prophylactically in the US, despite the lack of strong evidence. Often, parents receive little information about use and side effects of caffeine prenatally. There is variation in clinical indication for stopping caffeine and use of caffeine following discharge. Prospective studies should be aimed at determining optimal timing for initiation and discontinuation of caffeine, as well as to direct development of national guidelines.

INVASIVE VENTILATION DECREASES RENAL CAPILLARY SURFACE DENSITY MORE THAN NONINVASIVE RESPIRATORY SUPPORT IN TERM LAMBS

K. Albertine*, E Staub, M Dahl. University of Utah, Salt Lake City, UT

10.1136/jim-2019-WMRC.121

Purpose of study Ventilation, regardless of mode, decreases glomerular capillary surface density (SVgc) in the renal cortex of preterm lambs (Staub et al. 2017). We hypothesized that ventilation itself, not just renal immaturity, reduces SVgc. We compared SVgc of unventilated term lambs with two groups of ventilated term lambs: mechanical ventilation (MV) versus noninvasive respiratory support (NRS).

Abstracts
**Purpose of study** Hypoxia contributes to the development of pulmonary hypertension (PH) by promoting an uncoupled state of the enzyme endothelial nitric oxide synthase (eNOS) that produces superoxide instead of nitric oxide (NO). Folic acid has been shown to recouple eNOS and prevent the development of several hypoxia-induced markers of PH in a murine model. The impact of folic acid on recoupling eNOS in a piglet model of chronic hypoxia-induced PH is unknown. We tested the hypotheses that treatment with folic acid will increase eNOS recoupling in hypoxic piglet pulmonary artery endothelial cells (PAECs) and that treatment with folic acid will decrease pulmonary vascular resistance (PVR) in an in vivo piglet model of chronic hypoxia-induced PH.

**Methods** Piglet PAECs were cultured under hypoxic conditions (4% oxygen) for 48 hours with no folic acid or 1.0 μM folic acid. eNOS dimer-to-monomer ratios were measured by western blot. NO concentrations were quantified by chemiluminescence. Newborn piglets were obtained on DOL 2 and placed in a normobaric hypoxic chamber (10–11% oxygen) for 10 days. Some hypoxic piglets received 5 mg/kg/day of folic acid from days 3–10 of hypoxia while other hypoxic piglets received no folic acid. On DOL 10 hemodynamic measurements of pulmonary artery pressure, left ventricle diastolic pressure, and cardiac output were obtained to calculate PVR.

**Summary of results** Hypoxic PAECs treated with 1.0 μM folic acid (n=9) demonstrated increased eNOS dimer-monomer ratios compared to untreated hypoxic PAECs (n=9)(p<0.05). Hypoxic piglet PAECs treated with 1.0 μM folic acid (n=13) demonstrated a 16% increase in NO production compared to untreated hypoxic PAECs (n=13)(p<0.05). PVR was lower in hypoxic piglets treated with 5 mg/kg/day folic acid (n=5) compared to untreated hypoxic piglets (n=9)(p<0.05).

**Conclusions** Folic acid increases NO production by improving eNOS recoupling in hypoxic piglet PAECs. In an in vivo piglet model, treatment with folic acid ameliorates chronic hypoxia-induced PH. These findings suggest that folic acid has therapeutic potential to treat PH in human infants suffering from cardiopulmonary conditions associated with chronic hypoxia.

**FOLIC ACID TREATMENT RECOUPLES ENDOTHELIAL NITRIC OXIDE SYNTHASE AND AMELIORATES CHRONIC HYPOXIA-INDUCED PULMONARY HYPERTENSION IN NEWBORN PIGLETS**

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10.1136/jim-2019-WMRC.122
E Guslits*, M Steurer, H Nawaytou, R Keller. UCSF, San Francisco, CA

10.1136/jim-2019-WMRC.124

Purpose of study To evaluate B-type Natriuretic Peptide (BNP) as a longitudinal biomarker for clinical outcome in Congenital Diaphragmatic Hernia (CDH). We hypothesize that persistent elevation of BNP (>2 wks of age) is associated with mortality or need for prolonged respiratory support at 36 days (Steurer et al, 2014).

Methods used Retrospective cohort study. Detailed clinical information and BNP levels (pg/ml) were collected for the duration of hospitalization. At 36d, infants were classified as Poor Outcome if deceased or receiving ongoing respiratory support or Good Outcome if alive off support. Infants with congenital heart disease other than ASD, VSD, or PDA were excluded. BNP levels were measured at week 1 (n=35), 3 (n=43), 4 (n=37) and 5 (n=38); week 2 data were insufficient for analysis. At each time point, we generated receiver operator characteristic curves using log transformed BNP and defined the BNP cut-off that maximized correct classification.

Summary of results 22 VLBW infants with and 26 without PH were studied. Those with PH had a significantly longer birth weight. There were no other demographic differences between groups. The infants with PH had a significantly increased Rs and decreased Crs. Conclusions In this pilot study of VLBW infants screened for PH at 34–38 weeks PMA, those with PH had a significantly increased Rs and decreased Crs compared to those without PH. Additional studies are needed to further phenotype infants with evolving BPD and PH.

LONGITUDINAL B-TYPE NATRIURETIC PEPTIDE MEASUREMENT AS PROGNOSTIC MARKER FOR OUTCOMES IN CONGENITAL DIAPHRAGMATIC HERNIA

125 NONINVASIVE RESPIRATORY SUPPORT OF PRETERM LAMBS LEADS TO APPROPRIATE ALVEOLAR CAPILLARY GROWTH

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10.1136/jim-2019-WMRC.125

Purpose of study We reported that long-term outcomes for former-preterm lambs that were mechanically ventilated (MV) for 6–7d during the first week of postnatal life had persistent structural simplification of alveoli at 5 months corrected postnatal age (cPNA) compared to postnatal age-matched control term lambs. Five months cPNA is equivalent to ~6 yr PNA in humans (Dahl 2018). Most recently, we reported that noninvasive respiratory support (NRS) used for resuscitation and continuing management led to structural indices of alveolar formation that were comparable to the same indices in the lungs of control term lambs (Rebentisch, WSPR 2019; J Invest Med 67:91, 2019). However, the latter report did not assess alveolar capillary growth. Therefore, our aim was to quantify alveolar capillary growth in the lungs of the same former-preterm lambs that were managed by NRS and control term lambs.

Methods used Preterm lambs (~128d gestation; term ~150d; equivalent to lung development at ~28w gestation in humans) were resuscitated by NRS and continued with NRS for 6–7d (n=5; Rebentisch, J Invest Med 67:91, 2019). This NRS group was weaned from all respiratory support and lived for 5 months cPNA. Control term lambs lived for 5 months PNA herniation into the thorax was more common in Poor (90%) than Good (50%) Outcome (p=0.002). All infants with ECMO support had Poor Outcome (n=7/29, 24%; p=0.018). Median age at extubation was significantly longer in Poor vs. Good Outcome (19d, IQR 14–23 vs. 8d, IQR 4–11; p<0.001).

Median BNP at 1wk was lower in Poor Outcome (480, IQR 259–1020 vs. 1009, IQR 551–1540; p=0.08). At wks 3, 4 and 5, median BNP was higher in Poor Outcome: (3wks: 666, IQR 466–1040 vs. 111.5, IQR 76.5–227.5; p=0.0004); (4wks: 281, IQR 135–448 vs. 37.5, IQR 33–75.5; p=0.0018); and (5wks: 96, IQR 70–301 vs. 29, IQR 17–58; p=0.0025).

Conclusions Higher BNP at week 3–5 of life is associated with Poor Outcome in CDH. BNP cut-off points to predict Poor Outcome decline over this time frame.

Abstract 123 Table 1 Demographics and PFT results of infants without and with pulmonary hypertension (PH)

<table>
<thead>
<tr>
<th>Birth weight (g)</th>
<th>Gestational age (wks)</th>
<th>Male (%)</th>
<th>Caucasian (%)</th>
<th>PMA at time of PFT (wks)</th>
<th>Days of PFT from PH ECHO</th>
<th>Rs (cmH2O/ml/sec)</th>
<th>Crs (ml/cmH2O)</th>
<th>FRC/kg (ml/kg)</th>
<th>Number of Infants</th>
</tr>
</thead>
<tbody>
<tr>
<td>No PH (n=26)</td>
<td>Any PH (n=22)</td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
<td></td>
</tr>
<tr>
<td>945±181</td>
<td>823±224*</td>
<td>27.4±2.1</td>
<td>50</td>
<td>35.0±1.1</td>
<td>3.0±2.3</td>
<td>0.073±0.032</td>
<td>1.88±0.51</td>
<td>23.0±6.4</td>
<td></td>
</tr>
</tbody>
</table>

Abstract 124 Table 1 Area under the curve (AUC), and BNP cut-off values

<table>
<thead>
<tr>
<th>BNP Time Point</th>
<th>AUC (95%CI)</th>
<th>BNP cut-off (pg/ml, geometric mean)</th>
<th>Sensitivity</th>
<th>Specificity</th>
<th>Correctly Classified</th>
</tr>
</thead>
<tbody>
<tr>
<td>Week 1</td>
<td>0.67 (0.49–0.86)</td>
<td>539</td>
<td>83.3%</td>
<td>52.9%</td>
<td>68.6%</td>
</tr>
<tr>
<td>Week 3</td>
<td>0.82 (0.69–0.96)</td>
<td>285</td>
<td>88.9%</td>
<td>81.3%</td>
<td>86.1%</td>
</tr>
<tr>
<td>Week 4</td>
<td>0.82 (0.62–1)</td>
<td>101</td>
<td>88.0%</td>
<td>83.3%</td>
<td>86.5%</td>
</tr>
<tr>
<td>Week 5</td>
<td>0.82 (0.62–1)</td>
<td>61</td>
<td>85.2%</td>
<td>81.8%</td>
<td>84.2%</td>
</tr>
</tbody>
</table>

Values are Mean ± SD unless otherwise noted; FRC obtained in 19 versus 20 patients; * P criteria. Based on previous data, about 22 infants were needed per group to show a 25% difference in Rs with a α of 0.05 and power of 80%.

Summary of results 22 VLBW infants with and 26 without PH were studied. Those with PH had a significantly lower birth weight. There were no other demographic differences between groups. The infants with PH had a significantly increased Rs and decreased Crs.

Conclusions In this pilot study of VLBW infants screened for PH at 34–38 weeks PMA, those with PH had a significantly increased Rs and decreased Crs compared to those without PH. Additional studies are needed to further phenotype infants with evolving BPD and PH.
(equivalent to ~6 yr postnatal ages in humans). Quantitative histology was used to compare alveolar capillary surface density between the NRS group of preterm lambs and control term lambs.

Summary of results Former preterm lambs that were managed by NRS during the first week of postnatal life had the same alveolar capillary surface density (1,040±264 cm⁻¹; mean±SD; n=5) as control term lambs (918±61 cm⁻¹; n=5; not significant). Epithelial surface density, the reference space, was not different between the NRS group (4,274±843 cm⁻¹; n=5) and the control term group (3,938±447 cm⁻¹; n=5).

Conclusions NRS management, including resuscitation, of preterm lambs is associated with equivalent long-term growth of alveolar capillaries compared to control term lambs. These results, combined with our recent results, suggest that NRS management leads to appropriate capillary growth and structural formation of alveoli. Supported by R01 HL110002 and Division of Neonatology
normocapnia (temperature corrected PaCO₂38–48 mmHg) or hypercapnia (50–60 mmHg). Analysis was performed adjusting for hemoglobin (Hgb) level (high: 13–15 g/dL or low: 9–12 g/dL) and temperature.

Summary of results For pulmonary blood flow, high vs. low PaCO₂ resulted in a non-significant decrease in geometric mean blood flow of 3% (95% CI for relative change: 4 to 12%, p=0.41). For carotid blood flow (figure 1), high vs. low PaCO₂ resulted in a statistically significant increased geometric mean blood flow of 16% (95% CI for relative change: 0.1% to 34%, p<0.05).

Conclusions Within a clinically relevant PaCO₂ range of 38 to 60 mmHg, there is no significant alteration in pulmonary blood flow in lambs with HIE and PPHN. In contrast, hypercapnia is associated with an increase in carotid flow and possibly increased O₂ delivery to the brain. Clinical trials evaluating long-term effects of permissive hypercapnia (PaCO₂50–60 mmHg) are warranted in HIE with PPHN.

127 NASAL HIGH FREQUENCY JET VENTILATION VERSUS NASAL INTERMITTENT POSITIVE PRESSURE VENTILATION AS A MEAN OF POST EXTUBATION RESPIRATORY SUPPORT

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Purpose of study To describe our experience of successfully using nHFJV superimposed on to nasal intermittent positive pressure ventilation (NIPPV) after extubation or when NIPPV mode is failing to maintain blood gases in an acceptable range in ELBW infants.

Methods used All ELBW infants who received nHFJV while in the NICU were included in this study. IRB approval was obtained. Data was collected using NICU database and electronic medical records. The Bunnell Life Pulse HFJV was used to provide settings of: Jet PIP 30-35 cmH₂O, Jet Rate 240-300 bpm, Inspiratory time (IT) of 0.03 s. NIPPV rate on the conventional ventilator was set at 40 bpm, PIP at 30 cmH₂O, PEEP 8–10 cmH₂O, and IT at 0.5 s.

Summary of results Fourteen ELBW patients met the criteria for inclusion. Mean birth weight was 642 g and mean gestational age was 24.5 weeks. In 9 patients, nHFJV was used as a primary mode of extubation along with NIPPV after invasive HFJV treatment while in 5 patients it was used as a rescue mode when NIPPV settings were maximized with hypoxia and carbon dioxide retention. Twelve patients (86%) remained successfully extubated by 72 hours of treatment. Mean duration of invasive and non-invasive ventilation was 36 days and 45 days respectively. Duration of nHFJV was a mean of 9 days before discontinuing nHFJV. Twelve of 14 patients were on room air at the time of discharge.

Conclusions This is the first reported case series of the successful use of rescue nHFJV in conjunction with NIPPV to avoid reintubation in ELBW infants. Additional studies are needed to evaluate this mode of support in ELBW infants failing non-invasive modes of respiratory support.

128 NEUROBEHAVIORAL OUTCOMES OF FORMER PRETERM LAMBS ARE BETTER WHEN RESPIRATORY MANAGEMENT IS NONINVASIVE COMPARED TO INVASIVE

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Purpose of study Neurobehavioral impairments may be a long-term outcome for survivors of prematurity birth and invasive mechanical ventilation (MV). Former preterm (FPT) lambs develop diffuse brain damage, evident as increased apoptosis, and decreased proliferation, of neurons and glia compared to other premature lambs managed by noninvasive respiratory support (NRS) or term lambs that are not ventilated. We developed neurobehavioral tests of learning and memory for normal term lambs (Bowen, J Invest Med 64:294, 2016). We used these tests to assess the hypothesis that neurobehavioral outcomes will be better when management is by NRS compared to MV.

Methods used Three groups of FPT lambs were studied: (1) MV for 6–7d, n=6; (2) NRS, including for resuscitation, for 6–7d, n=5; and (3) MV for ~3h followed by NRS for 6–7d, n=5. The FPT lambs lived for 6 months (3 months corrected postnatal age, cPNA; equivalent to ~6 years for humans). Control term lambs were the fourth group (n=8). Tests were repeated at 2 and 5 months. Lambs were habituated to the tests before having two 10-min trials/test (one test/d). Tests were separated by 30-min of rest. The routine was repeated the next day. Different tests were separated by a rest day. One test used a mirror and a non-reflective surface to test socialization. Another test used a maze, for which the reward was eew’s milk, to test memory and problem solving.

Summary of results For the 5-month test, 6–7d MV FPT lambs spent significantly more time at the non-reflective surface (8 (24) sec; median (IQR) than NRS FPT lambs (1 (8) sec; p<0.05). No differences were detected between the NRS FPT, MV for ~3h then NRS FPT, and control term groups. For the 5-month maze test, no differences in time to find the reward were detected between the 6–7d MV FPT lambs (7 (134) sec) and NRS FPT group (5 (2) sec).

Conclusions Our results suggest that FPT lambs that were managed by MV during the first week of postnatal life have worse long-term neurobehavioral outcomes than NRS FPT lambs. The range for the MV FPT lambs implies variability that may provide opportunity to identify underlying mechanisms. (HL110002 and Division of Neonatology).

129 EXPOSURE TO CANNABIDIOL (CBD) RESULTS IN MYOCGENIC DIFFERENTIATION AND ALTERED FUNCTION IN FETAL RAT LUNG FIBROBLASTS

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Purpose of study Previously, we have demonstrated that perinatal nicotine exposure disrupts specific epithelial-mesenchymal paracrine signaling pathways, resulting in pulmonary interstitial fibroblast transdifferentiation to a myogenic phenotype.
Abstracts

Recently, cannabidiol (CBD) has been implicated as one of the emerging contributors to the recent teen lung disease crisis in the US. However, it is unknown how CBD affects lung fibroblast differentiation and the developing lung.

Since a balance between Wnt and PPARγ signaling is a key determinant of lung interstitial fibroblast phenotype, i.e., lipogenic vs. myogenic, using a well-established in vitro model, we tested the hypothesis that CBD exposure would up-regulate Wnt and down-regulate PPARγ pathways in fetal rat lung fibroblasts.

Methods used Using established methods, embryonic day 19 Sprague-Dawley rat lung fibroblasts were treated for 24 or 72 h with CBD (10⁻⁴, 10⁻⁵, 10⁻⁶ M). Rate of triglyceride uptake, a key indicator of lipofibroblast function, was assessed through [³H]-triolein uptake assay. Mesenchymal markers of differentiation were determined by western analysis and qRT-PCR.

Summary of results As assessed by differentiation and functional assays, CBD treatment affected fibroblast phenotype dose- and time-dependently. However, contrary to our hypothesis, both lipogenic and myogenic markers increased significantly at both time-points examined with more profound changes following 72 h exposure. In particular, Wnt signaling/myogenic markers (LEF-1, β-catenin, Fibronectin, Calponin, and α-SMA) demonstrated more robust changes, indicating CBD-induced myogenic differentiation of pulmonary interstitial fibroblasts, which was corroborated by triolein uptake assay (p<0.05).

Conclusions We conclude that in line with the previously known effects of nicotine on the developing lung fibroblasts, CBD also induces myogenesis. However, unlike nicotine’s effects on lung fibroblasts, CBD’s effects seem to be more complex, i.e., concomitant up-regulation of both lipogenic and myogenic markers, possibly reflecting a compensatory response to CBD-induced dyshomoeostasis. The implications of our data in an incisor system are unclear and are being studied using well-established in vivo rodent models.

EVIDENCE FOR PULMONARY GROUP 2 INNATE LYMPHOID TYPE II CELL INVOLVEMENT IN PERINATAL NICOTINE-INDUCED ASTHMA

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Purpose of study Exposure of the developing lung to nicotine is a significant contributor to offspring asthma and other allergic diseases. Though the mechanisms underlying perinatal nicotine-mediated pulmonary structural effects are well studied, the mechanisms underlying allergic predisposition remain unknown. Group 2 innate lymphoid type II cells (ILC2s), relatively new players in asthma pathogenesis, produce Th2 cytokines in response to a number of environmental signals; however, their involvement in perinatal nicotine-induced asthma is unknown. We tested the hypothesis that ILC2s are involved in perinatal nicotine-induced asthma.

Methods used Using a well-established perinatal nicotine exposure mouse model, pregnant C57BL/6J mouse dams received nicotine (2 mg/kg, s.c.) or diluent once daily from e6 to postnatal day (PND) 21. Starting 3 days before sacrifice on PND21, pups were administered recombinant murine (rm) IL-33 (0.5μg intranasally) or PBS once daily under brief isoflurane anesthesia, and sacrificed 24h after the last rmIL-33 dose for ILC2s collection using Aria III cell sorter. ILC2s were characterized based on the lack (CD3−, CD45R−, Gr-1−, CD11c−, CD11b−, Ter119−, TCRγδ−, and FcγRI−) and the expression of (CD45, ST2, and CD117) classical lineage markers. Cultured (5 × 10⁴ cells/well) ILC2s from control and nicotine exposed pups were treated with rmIL-33 (10 ng/ml) for 48h and the supernatant was analyzed for Th2 cytokines IL-5 and IL-13. Lastly, the effect short chain fatty acid butyrate on IL-33-induced production of Th2 cytokines by ILC2s was determined.

Summary of results Nicotine treatment resulted in a greater yields of ILC2s (ILC2s/lung/mouse- 2770±72 vs 1790±89; nicotine vs control; N=4; p<0.001). IL-33 treatment of nicotine exposed ILC2s resulted in a higher Th2 cytokine production compared with ILC2s isolated from controls (N=4; p<0.001). Sodium butyrate (20 mM) decreased IL-5 production by 61% and IL-13 production by 89% (vs. IL-33 treated group).

Conclusions We for the first time implicate the involvement of ILC2s in perinatal nicotine-induced asthma and suggest a novel novel, safer, and effective mechanism-based (ILC2 negative regulators) therapy for perinatal smoke/nicotine-induced asthma.

COMMON VAPE FLAVORS ARE DETERIMENTAL TO LUNG FIBROBLAST DIFFERENTIATION

Y Liu, R Sakurai, Y Wang, A Harb*, V Rehan. Lundquist Institute for Biomedical Innovation at Harbor-UCLA Medical Center, Torrance, CA

Purpose of study A significant strategy of electronic cigarette (e-cig) marketing focuses on the variety of vape flavors available, which often mimic the taste of fruits to more elaborate aromas. Manufacturers of e-liquids, the fluid used in e-cigs, argue that their flavored products are ‘food grade’ and are ‘generally recognized as safe’ (GRAS). However, the GRAS certification pertains only to ingestion and does not take into account inhalation nor the heat-induced changes in flavorings’ compositions of vapes. The flavoring chemical concentrations, 1–4% range (10–40 mg/mL) in e-liquids are high enough to raise concern for toxicity, particularly, with chronic exposure. Since interstitial fibroblasts are the primary determinants of lung phenotype, we examined the effect of select commonly used e-cig flavorings on cultured fetal rat lung fibroblast toxicity and differentiation.

Methods used Using well established methods, embryonic day 19 Sprague-Dawley rat lung fibroblasts were isolated and were treated for 24 h with nicotine (1 × 10⁻⁶M), vanilla, cherry, and menthol (1 × 10⁻⁹ and 1 × 10⁻⁶M) either alone or nicotine + each flavoring combination. Fibroblast toxicity [lactate dehydrogenase assay (LDH)] and differentiation (protein levels of key lipogenic (PPARγ) and myogenic proteins (α-SMA and fibronectin) via western analysis and immunofluorescence staining) were determined.

Summary of results Based on the LDH assay, there was no overt effect on fibroblast toxicity among any of the flavorings examined at either tested doses (1 × 10⁻⁹ and 1 × 10⁻⁶M). However, with the exception of menthol and menthol+nicotine, there were marked effects on fibroblast differentiation
in all other groups as evidenced by significantly decreased PPARγ and increased α-SMA and fibronectin protein levels as well as positive staining for these markers.

**Conclusions** Based on our data, common e-liquid flavorings demonstrated significantly detrimental, albeit, differential effects on lung fibroblast differentiation as demonstrated by reduced lipogenic but increased myogenic markers, which are key characteristics of myofibroblasts. These findings, in general, contradict the claims of e-liquid manufacturers. However, more extensive and in vivo studies need to be performed to determine the clinical significance of our data.

**Neuroscience I**

**Concurrent session**

3:15 PM

Thursday, January 23, 2020

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### Abstract 132

**DURATION OF AMERICAN FOOTBALL PLAY AND CHRONIC TRAUMATIC ENCEPHALOPATHY**

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10.1136/jim-2019-WMRC.132

**Purpose of study** Chronic traumatic encephalopathy (CTE) is a neurodegenerative disease associated with exposure to contact and collision sports, including American football. We hypothesized a dose-response relationship between duration of football played and CTE risk and severity.

**Methods used** In a convenience sample of 266 deceased American football players from the Veterans Affairs-Boston University-Concussion Legacy Foundation and Framingham Heart Study Brain Banks, we estimated the association of years of football played with CTE pathological status and severity. We evaluated the ability of years played to classify CTE status using receiver operating characteristic curve analysis. Simulation analyses quantified conditions that might lead to selection bias.

**Summary of results** 223 of 266 participants met neuropathological diagnostic criteria for CTE. More years of football played was associated with having CTE (odds ratio [OR] = 1.30 per year played, 95% CI, 1.19–1.41; P=3.8 × 10⁻⁵) and with CTE severity (severe vs. mild; OR=1.14 per year played, 95% CI, 1.07–1.22; P=3.1 × 10⁻⁵). Participants with CTE were 1/10th as likely to have played <4.5 years (negative likelihood ratio [LR]=0.102, 95% CI, 0.100–0.105) and were 10X as likely to have played >14.5 years (positive LR=10.2, 95% CI, 9.8–10.7) compared with participants without CTE. Sensitivity and specificity were maximized at 11 years played. Simulation demonstrated that years played remained adversely associated with CTE status when years played and CTE status were both related to brain bank selection across widely ranging scenarios.

**Conclusions** The odds of CTE double every 2.6 years of football played. After accounting for brain bank selection, the magnitude of the relationship between years played and CTE status remained consistent.

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### Abstract 133

**LONGITUDINAL FUNCTIONAL MAGNETIC RESONANCE IMAGING (fMRI) FINDINGS IN ADOLESCENTS**

1,2K Nikroo*, 3A Hawkins, 2D Singhania, 2S Chitoori, 2P Wiederkehr, 2P Otero, 2,3B Afghani. 1University of Southern California, Los Angeles, CA; 2UC Irvine School of Medicine, Irvine, CA; and 3CHOC Hospital, Orange, CA

10.1136/jim-2019-WMRC.133

**Purpose of study** Studies that have evaluated functional MRI in adolescent cannabis (marijuana) users at a single time point have shown alteration in brain connectivity, especially in the orbitofrontal/temporal areas. However, longitudinal studies of fMRI at two different time points is limited. The objective of this study was to evaluate the fMRI changes in adolescent cannabis users over time.

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### Table 1

**Longitudinal MRI Findings in Cannabis (Marijuana) Users**

<table>
<thead>
<tr>
<th>First Author, Year Published, Location</th>
<th>Number of subjects vs. controls and age at initial MRI</th>
<th>Time period to follow-up MRI</th>
<th>Frequency of marijuana (MJ) use</th>
<th>fMRI findings in subjects compared to controls</th>
</tr>
</thead>
<tbody>
<tr>
<td>Cheetham, 2012, Australia</td>
<td>Subjects: N=30, Heavy MJ and Alcohol users Controls: Non-users, N=38, age at baseline for both groups=16–19 yr</td>
<td>4 years from baseline</td>
<td>Subjects: median=16.6 times</td>
<td>Smaller (thinner) orbitofrontal cortex volumes at age 12 years but thicker cortices at f/u in cannabis users compared to non-users (p=0.05)</td>
</tr>
<tr>
<td>Jacobus, 2015, USA</td>
<td>Subjects: Heavy MJ, N=22, Controls: Non-users, N=43, age at baseline for both groups=16–19 yr</td>
<td>1.5 and 3 years from baseline</td>
<td>Subjects: &gt;100 lifetime episodes Controls:</td>
<td>Marijuana and alcohol showed thicker cortical estimates across the brain (23 regions), particularly in frontal and parietal lobes (p&lt;0.05)</td>
</tr>
<tr>
<td>Camchong, 2016, USA</td>
<td>Subjects: MJ users, N=22, age at baseline=13–23 yrs Non-users: N=63, age at baseline=10–22 yrs</td>
<td>1.5 years from baseline</td>
<td>Subjects: means=1049 at baseline and 1202 days during flu period, Controls: 0 days to 22 days during flu period</td>
<td>Cannabis users showed a decrease in functional connectivity between caudal anterior cingulate cortex and dorsolateral and orbitofrontal cortices across time (p&lt;0.05)</td>
</tr>
<tr>
<td>Tervo-Clemmens, 2018, USA</td>
<td>Subjects: MJ users, N=22 Non-users: N=63, age at baseline for both groups=12 yrs</td>
<td>3 years from baseline</td>
<td>Subjects: # Joints/day between 13–15 years (follow-up period)=0.076 Non-users=0</td>
<td>Those who would initiate cannabis use by 15 years of age had activation differences in frontoparietal (increased) and visual association (decreased) regions and poorer executive planning scores (Stockings of Cambridge) compared with noninitiators (p&lt;0.05)</td>
</tr>
</tbody>
</table>
Methods used A comprehensive literature review was performed through multiple search engines, such as PubMed, Google Scholar, and Cochrane using keywords, ‘fMRI’, ‘cannabis’, ‘marijuana’, and ‘longitudinal’. Only studies that involved 2 or more fMRIs at least 18 months apart in adolescent cannabis users and a non-user group were included in our analysis.

Summary of results Of 20 studies, 4 satisfied our inclusion criteria (table 1). Adolescent cannabis users had thinner cortices pre-initiation of cannabis but compared to non-users thicker cortices at follow-up (after cannabis use). Although cause and effect can’t be proven, thinner brain cortex in frontal and orbitofrontal volume may predict cannabis use later in life. In addition, there was a decrease in functional connectivity and impact on executive planning in cannabis users over time. The studies accounted for some, but not all of the confounding variable. There was variation in chronicity, age at onset, and amount of cannabis used.

Conclusions Longitudinal fMRI studies in cannabis users are limited but there is some evidence that cannabis causes alterations in certain areas of the brain that may lead to deleterious effects. However, it is not clear if the alteration is due to use of cannabis or other confounding variables such as stage of adolescent maturation, use of other substances or preexisting differences. Prospective, longitudinal studies are needed to evaluate the association between chronicity of cannabis use and fMRI findings, taking into account other confounding variables.

### Abstract 134

**BDNF AUGMENTATION BY AMPAKINES EXPRESSION RESCUES SYNAPTIC DYSFUNCTION AFTER GLOBAL CEREBRAL ISCHEMIA IN THE DEVELOPING BRAIN**

N Chalmers, JE Orfila, PS Herson, RM Dietz*. Univ of Colorado, Aurora, CO

10.1136/jim-2019-WMRC.134

**Purpose of study** Global ischemia in the developing brain often leads to poor neurologic outcomes, including learning and memory deficits. Using a novel murine model of juvenile cardiac arrest (CA), we investigated mechanisms of cognitive deficits and recovery. Brain-derived neurotrophic factor (BDNF)-tyrosine kinase (TrkB) B signaling is a required pathway for memory formation and hippocampal synaptic plasticity. We recently demonstrated that enhanced BDNF-TrkB signaling contributes to recovery of cognitive function after global ischemia. AMPA/kines are allosteric modulators of AMPA receptors and some have been shown to augment BDNF levels. We hypothesize AMPAkines that augment BDNF will reverse synaptic dysfunction following CA.

**Methods used** Male juvenile mice (PND 20–25; equivalent to 2–3 year old human) were subjected to 8 min CA and resuscitated. Memory function was measured using contextual fear conditioning, a hippocampal-dependent memory task. Hippocampal CA1 long-term potentiation (LTP), a well-accepted cellular model for learning and memory, was measured in acute brain slices following a theta-burst stimulation (TBS, 40 pulses 100Hz). Increase in field excitatory post-synaptic potential (fEPSP) slope 60 min after TBS was analyzed to quantify LTP. BDNF ELISA from hippocampal tissue was performed per manufacturer’s instructions. Results reported as mean±SD.

**Summary of results** Memory is impaired 7d after CA (52±15% sham, n=8, vs 24±19% CA, n=9, p<0.05) and recovers 30d after CA (52±14%, n=7). Hippocampal LTP correlates with this neurobehavioral recovery. In controls, LTP was 153±15% (n=8) of baseline (100%). In contrast, 7d after CA, LTP was impaired (110±16%, n=8, p<0.05 vs sham), followed by recovery at 30d (157±16%, n=8, p<0.05). Using ELISA, BDNF levels decreased after CA compared to sham animals (n=4, p<0.05). Injection of the BDNF-inducing AMPAkine LY404187 7d after CA or sham increased BDNF (n=4 each, p<0.05). In paired studies 7 days after CA, LY404187 applied to hippocampal slices 7d after CA rescues impaired LTP (111±12 [n=4] vs. 153±13 [n=4], p<0.05, paired t-test).

**Conclusions** These data provide unique new data that BDNF enhancing drugs are capable of enhancing recovery from global cerebral ischemia in the young brain during critical school ages.

### Abstract 135

**EFFECTS OF GSK3B & MTOR ON MICROTUBULE STABILITY & NEURITE LENGTH**

1. H Kornblum, K Ludwig, S Ravain. ¹UCLA, Los Angeles, CA; ²Tulane University, New Orleans, LA

10.1136/jim-2019-WMRC.135

**Purpose of study** In our study, we hypothesized that modifying GSK3B or mTOR will ultimately result in increased microtubule stability and increased neurite length, which could lend insights into potential avenues for treating glioblastoma (GBM). Therefore, we investigated whether altering GSK3B or mTOR could promote microtubule stability and increase neurite length in mouse neuronal cells in vitro.

**Methods used** Mouse neurons were treated with DMSO (control and a strong dissociator), CHIR (inhibits GSK3B), Rapa (inhibits mTOR), Epothilone (stabilizes microtubules), and ST401 (destabilizes microtubules). The mouse neurons were derived from mouse stem cells. The differentiation process was straightforward. We took mouse neurons, plated them, and then withdrew growth factors from their media and waited approximately one week.

**Summary of results** The average neurite lengths of mouse neurons placed in DMSO was roughly 174um, 347um in EPO, and 341um in Rapa. This shows that Rapa had the most stabilizing effect on the microtubules, followed by EPO and DMSO. CHIR had little to no effect on microtubule stability.

**Conclusions** These data provide new insights into potential avenues for treating glioblastoma (GBM) by investigating the effects of GSK3B and mTOR on microtubule stability and neurite length in mouse neuronal cells in vitro.
177 μm in ST, 257 μm in CHIR, and 141 μm in rapamycin. We observed that the mouse neurons treated with EPO and CHIR appeared more pronounced, clear, and longer, compared to those treated with rapamycin and ST. 

**Conclusions** We concluded that inhibiting GSK3β promotes microtubule stability and increases neurite length, and inhibiting mTOR promotes microtubule stability, but decreases neurite length. Future directions include evaluating the molecular response and resistance to mTOR inhibition to improve options for GBM treatment by exploiting vulnerabilities in mTOR targeting and microtubule stabilization.

**Summary of results** Plk1 and Nusap1 expression were restricted to a subset of olig2 cells located in the progenitor domain, consistent with the possibility that they mark progenitors fate-restricted to specify into oligodendrocyte-lineage cells. In addition, outside the progenitor domain, plk1 was colocalized with olig2 positive cells but there was little colocalization between nusap1 and olig2. Based on the experimentation on mutant zebrafish, oligodendrocyte development seems to be delayed in the absence of Boc receptors for Sonic Hedgehog (Shh) and premature in the absence of Fbxw7-dependent degradation of Notch. 

**Conclusions** The results provide insight into how the Shh and Notch signalling pathways regulate oligodendrocyte development. In addition, our work supports the theory that a fate-restricted progenitor cell differentiates into oligodendrocyte-lineage cells. We suggest that colocalization between only plk1 and olig2 marks oligodendrocyte precursor cells. Furthermore, our work indicates that inducing the expression of plk1, nusap1, and olig2 in pluripotent neural stem cells can ultimately create oligodendrocytes that remyelinate damaged axons and restore normal function. Therefore, this research has significance in helping to create a therapy for patients whose myelin has been damaged by disorders such as multiple sclerosis.
morpholino oligonucleotide conjugated with a cell penetrating peptide reduced glycogen, however was nephrotoxic. Antisense Oligonucleotides (ASO) technology has emerged as a powerful therapeutic alternative for the treatment of genetic disorders by targeting RNA. Most recently, therapy for spinal muscular atrophy has been successful using ASOs, and our hope is that ASO technology will be successful in Pompe disease. We propose that knocking down GYS1 with ASOs may prove to be a useful therapeutic target in Pompe disease.

**Methods used** In order to impart specificity for the muscle glycogen synthase (GYS1), we used ASO-mediated gene silencing through the RNaseH1 dependent degradation mechanism.

**Summary of results** Over 150 ASOs were designed and screened in vitro to identify the most efficacious ASO for testing in wild type mice. The lead from the screen were validated in a dose response study and the top 10 ASOs were screened in vivo. Three ASOs (GYS1 ASO#1, ASO#2 and ASO#3) showed the best tolerability and efficacy profile leading to knock down of GYS1 mRNA by approximately 50% of control. We performed a pilot study of the efficacy of three GYS1 ASOs in Pompe mice as monotherapy. We were able to see reduced muscle GYS1 mRNA levels and glycogen with the ASOs versus PBS or a mismatch ASO. ASO#2 however resulted in weight loss of the mice. Overall GYS1 ASOs #1 and possibly #3 seemed the most promising in reducing muscle GYS1 levels and glycogen content.

**Conclusions** These preliminary studies provide proof of principle that GYS ASOs might be a potentially promising adjunct treatment for Pompe disease in reducing GYS1 and glycogen in muscle.

139  **PRESUMED CONVERSION DISORDER IN A PATIENT FOUND TO HAVE NEW ONSET MULTIPLE SCLEROSIS: CHALLENGES IN DIAGNOSIS AND TREATMENT**

1,5 Lam*, 1,2E Laboy-Gonzalez, 1,3H Kotayukio, 1'Thara, 1'Abdijadid. 1Kern Medical Center, Bakersfield, CA; 2Ross University School of Medicine, Bridgetown, Barbados; 3American College of Physicians

**Introduction** Conversion Disorder, referred to as Functional Neurological Symptom Disorder under DSM-5, is often a diagnosis of exclusion after primary organic/neurological causes are ruled out, and when clinical findings do not correlate with recognized disease states. The following case report highlights the diagnostic and treatment challenges in a patient initially thought to have conversion disorder but later diagnosed with severe relapsing-remitting multiple sclerosis (MS).

**Case report** 45-year-old Caucasian female with a history of schizoaffective disorder and mild cognitive delay who was admitted to our facility after reported inability to move her body. She was then given fluoxetine in their drinking water or plain drinking water. The mice were sacrificed five weeks post-surgery and their brains harvested for Western blot analyses of Ki-67 (marker for neuron proliferation), doublecortin (DCX, protein expressed in immature neurons and neurons of the AHN), whereas selective serotonin reuptake inhibitors (SSRIs) stimulate it. Therefore, SSRIs may be a potential therapeutic approach to preventing and/or treating disruption of AHN and the subsequent CI. This investigation sought to test the hypothesis that the SSRI, fluoxetine, can reduce or block androgen deprivation-induced reduction of AHN.

**Methods used** Mice were randomly divided into four groups: sham/vehicle, sham/fluoxetine, castration/vehicle, and castration/fluoxetine. They underwent castration or sham surgery and were then given fluoxetine in their drinking water or plain drinking water. The mice were sacrificed five weeks post-surgery and their brains harvested for Western blot analyses of Ki-67 (marker for neuron proliferation), doublecortin (DCX, protein expressed in immature neurons and neurons of the AHN), whereas selective serotonin reuptake inhibitors (SSRIs) stimulate it. Therefore, SSRIs may be a potential therapeutic approach to preventing and/or treating disruption of AHN and the subsequent CI. This investigation sought to test the hypothesis that the SSRI, fluoxetine, can reduce or block androgen deprivation-induced reduction of AHN.

**Summary of results** Our investigation showed increased Ki-67 protein levels in castration/vehicle compared to the sham/vehicle group. The decrease in Ki-67 in the castration/vehicle groups was prevented in the castration/fluoxetine group. Furthermore, there were no differences between sham/vehicle and castration/fluoxetine groups. There were no differences in NeuN and DCX levels among the four treatment groups.

**Conclusions** The results suggest that treatment with fluoxetine reduced the effects of ADT on AHN. It will be important to test whether other drugs that stimulate AHN also are effective in reducing the effects of ADT and whether they can reverse the effects once they develop. This approach may be useful in preventing and/or treating ADT-induced CI in patients with prostate cancer.
Purpose of study Data are limited regarding current practice and outcomes for emergency department airway management in asthma. Our goal was to describe the methods, equipment used, medications, techniques, adverse events and outcomes for emergency airway management in asthmatic patients through multi-center surveillance.

Methods used We analyzed all intubations with a primary indication of asthma over a three-year period (January 1, 2016–December 31, 2018) using the National Emergency Airway Registry (NEAR), a 25-center prospective observational registry of ED intubations. We report the incidence of intubations for asthma, methods and medications used, device, peri-intubation adverse events, as well as intubation success and failure using univariate descriptive statistics and cluster-adjusted 95% Confidence Intervals (CI).

Summary of results A total of 19,071 encounters were recorded during the study period with 14,517 intubated for medical indications. Of those, 173 (1.2%, CI 0.9–1.6%) were intubated for asthma. Nearly two-thirds (n=56, 62.9%) were pre-oxygenated with bi-level positive airway pressure. Rapid sequence intubation (RSI) was used in 96.5% of cases. For these, more than half (n=85, 50.9%, CI 30.1–71.4%) were induced with ketamine, followed by etomidate at 48.5% (n=81, CI 27.8–69.7%). There was an overall adverse event rate of 12.14%. First attempt success was 90.8%. Overall intubation success was 100%. There was no difference in first-attempt success between ketamine and etomidate intubations. Compared to the registry as a whole, asthma patients were more likely to undergo RSI (96.5% versus 80.8%), pre-oxygenation with bi-level positive airway pressure (62.9% versus 6.2%), and induction with ketamine (50.9% versus 12.8%).

Conclusions In our registry, intubation for asthma is uncommon. When required, the majority of patients are intubated using rapid sequence intubation after pre-oxygenation with bi-level positive airway pressure and induction with ketamine, a practice unique to asthma. It is unclear whether the high rate of Bi-PAP use is reflective of initial treatment or simply the most popular means of pre-oxygenation. Videolaryngoscopy use and first-attempt success are high and similar to the registry as a whole.

Pulmonary and critical care I

Concurrent session

3:15 PM

Thursday, January 23, 2020

142 EMERGENCY DEPARTMENT AIRWAY MANAGEMENT FOR ASTHMA

1JF Godwin*, 2JF Riker, 3R Baker, 4J Madisen, 5RM Walls, 6C Brown, 7University of Utah, School of Medicine, Salt Lake City, UT; 8Bingham and Women’s Hospital, Boston, MA

Purpose of study Data are limited regarding current practice and outcomes for emergency department airway management in asthma. Our goal was to describe the methods, equipment used, medications, techniques, adverse events and outcomes for emergency airway management in asthmatic patients through multi-center surveillance.

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mortality data were extracted from the electronic medical record. We examined the association between delirium and depressive symptoms/HRQL with linear mixed effects models and the association between delirium and mortality with cox proportional hazard models. Models were adjusted for age, sex, native disease, and lung allocation score (LAS).

Summary of results ICU delirium occurred in 34 subjects (14%). Delirium was associated with worse generic physical HRQL but not significantly poorer depressive symptoms (table 1). Those who experienced delirium had a nearly 16-fold increased risk of mortality (table 2).

Abstract 143 Table 1 Differences in patient reported outcome scores by delirium status after LT

<table>
<thead>
<tr>
<th>Outcome Measure</th>
<th>Difference between those who were delirious and those who were not</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>SF12-PCS</td>
<td>4.0 (95% CI: -75, -65) [MCID: 5]</td>
<td>0.02</td>
</tr>
<tr>
<td>GDS</td>
<td>0.4 (95% CI: -0.7, 1.5) [MCID: 1.7]</td>
<td>0.49</td>
</tr>
</tbody>
</table>

MCID = Minimally clinically important difference

Abstract 143 Table 2 The association between delirium and risk of death through first post-transplant year

<table>
<thead>
<tr>
<th>Model</th>
<th>Hazard ratio (95% CI)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Unadjusted</td>
<td>11.3 (3.3, 38.4)</td>
<td>0.0001</td>
</tr>
<tr>
<td>Adjusted for age, sex, native disease, LAS</td>
<td>15.9 (3.9, 64.4)</td>
<td>0.0001</td>
</tr>
</tbody>
</table>

Conclusions Delirium after LT identifies a group at high risk for worse outcomes and is associated with poorer HRQL and increased mortality. Implementing strategies to prevent and/or treat delirium may improve both HRQL and mortality risk after LT.

Management of Severe Pulmonary Hypertension in Pregnancy with Epoprostenol

Case Report Patient was started on intravenous Epoprostenol in the intensive care unit at 27 weeks and 5 days gestation. The initial dose was 2ng/kg/min with increasing dose by increments of 1–2ng/kg/min every fifteen minutes until the side effects of flushing, jaw pain, nausea, or hypotension occurred. Epoprostenol discontinued due to extreme lethargy. Patient continued to endorse shortness of breath and lethargy, requiring transfer to a higher level of care. After arrival and prior to Epoprostenol being restarted, the fetal heart tracing showed decelerations prompting emergent cesarean at 28 weeks gestation. Postpartum, patient underwent second course of treatment with Epoprostenol and discharged with Sildenafil and Macitentan. Postpartum echocardiography did not demonstrate improvement or worsening of PAH. Within the first postpartum week patient experienced multiple pulmonary embolisms requiring treatment with heparin later transitioning to apixaban. The neonate was delivered via cesarean and transferred to NICU.

Therapeutic agents include endothelin receptor antagonists, phosphodiesterase-5 inhibitors, and prostacyclin analogs. Based on limited data, risk of fetal harm and maternal side effects remains uncertain. There are few case reports describing successful treatment with Epoprostenol in pregnant women with PAH. Endothelin receptor antagonists not used due to teratogenic effects. Further investigation is needed to formulate a safe and efficacious treatment plan.

Worse Sleep Quality is Associated with Increased Risk of Chronic Obstructive Pulmonary Disease Exacerbation

Purpose of study Sleep disturbance is recognized as a symptom component in chronic obstructive pulmonary disease.[i] One study demonstrated that poor sleep quality as defined by Pittsburg Sleep Quality Index (PSQI) score >5 is associated with increased risk of COPD exacerbations.[ii] However, limited sample sizes and confounders have impeded the ability to replicate or quantify this association.

Methods used A cross-sectional study of 1,137 participants with COPD from the Subpopulations and Intermediate Outcome Measures in COPD Study (SPIROMICS) were included for this analysis. SPIROMICS is a multi-center prospective cohort of subjects with and without COPD. The Pittsburgh Sleep Quality Index (PSQI) is an instrument with 19 scored responses to assess sleep quality. Participants with an index of 5 or higher were considered to have poor sleep. Exacerbations were defined as events that led to some contact with the healthcare system with use of systemic corticosteroids or antibiotics. We used negative binomial regression to examine the association between COPD exacerbations and the PSQI. The model was adjusted for age, gender, race, FEV1%predicted, GOLD severity, diagnosis of asthma, ischemic heart disease, apnea, depressive and anxiety symptoms, pneumococcal and influenza vaccination status, history of malignancy, income, and educational attainment.

Summary of results 609 (53%) subjects had poor sleep. Compared with those with good sleep quality, poor sleepers were...
RECURRENT SYNCOPE AS A PRESENTATION OF PULMONARY EMBOLISM – A CASE REPORT

K Suwanwongse,* N Shabaneh. Lincoln Medical and Mental Health Center, Bronx, NY

10.1136/jim-2019-WMRC.146

Case Report The diagnosis of pulmonary embolism (PE) is challenging especially when patients present with vague symptoms. Early diagnosis and management of PE is crucial to prevent life-threatening sequelae.

A 60-year-old woman bought by EMS after having syncope while walking. She reported shortness of breath (SOB) before syncope. She has hypertension, type 2 DM, osteoporosis and multiple syncopes (>10) in past year. Her investigation in another hospital includes EKG, 24-hour telemetry, echocardiogram and stress-test were normal but did not have chest CT. Her vital signs and exam were normal. Blood tests and EKG were unremarkable except mild anemia (Hb10.9) and elevation of troponin T (0.015). She was advised to admit but signed out against advice. On way out, she had another syncope that recovered spontaneously but had worsening SOB. She had tachypnea, respiratory distress and hypoxemia, oxygen saturation in 60s. Chest CTA showed prominent bilateral PE. She received heparin and thrombolysis (EKOS). Doppler sonography found occlusive thrombosis left popliteal vein with partially occlusive distal left superficial femoral vein. She was discharged on apixaban. Her SOB improved and no recurrent syncope was reported.

PE is a differential diagnosis for syncope in most textbooks, but when patients came due to syncope, PE, a potentially fatal disease requiring urgent attention, is rarely considered. This case is interesting as a patient had multiple syncopes without suspicious of PE until developing severe hypoxemia. PE should be considered in all patients experiencing syncope particularly with SOB, respiratory distress and hypoxemia.

COMPARISON OF LUNG INFLAMMATION IN A HOUSE DUST MITE- AND ASPERGILLUS FUMIGATUS EXTRACT-INDUCED ALLERGIC ASTHMA MOUSE MODEL

1) J Sumsion*, 1A Pulipher, 1C Pollard, 1A Benally, 1J Alt. 1University of Utah School of Medicine, Salt Lake City, UT; 2Fort Lewis College, Durango, CO

Purpose of study Asthma affects 8.3% of the U.S. population with rising prevalence and no cure. Current investigations have demonstrated that asthma is multifactorial. Due to these etiological complexities, animal models have been developed to further our understanding of the underlying pathogenesis and pathophysiology of different asthma phenotypes. The most well-characterized mouse model of allergic asthma is the ovalbumin model. However, asthma models induced by house dust mite (HDM) invasion have been recently considered to be more relevant to human asthma. We propose a new Aspergillus fumigatus (AF) extract-induced asthma model that may also be relevant to human asthma. Herein, we characterized and compared lung inflammatory responses induced by HDMs and AF extracts in two different mouse models of asthma.

Methods used BALB/c mice were challenged with (1) intraperitoneal and intranasal HDMs over 16 days or (2) three times weekly intranasal AF extracts for 8 weeks. Bronchoalveolar lavage (BAL) fluid samples were collected and subjected to differential leukocyte analysis using flow cytometry. Data are represented as the mean of detected CD45+ macrophages, CD11b+ lymphocytes, Ly-6G+ neutrophils and Siglec F+/MHCIIC+ eosinophils as percentages of total counted viable leukocytes.

Summary of results The models produced unique distributions of leukocytes in the BAL fluid. While the control group was almost exclusively characterized by CD11c+ alveolar macrophages (99.95%), the HDM-induced disease group demonstrated a significant neutrophil (39.06%) response. Contrastingly, the AF extract-induced disease group had significant increases in eosinophil (28.51%) and lymphocyte (14.52%) populations compared to controls (0.37% and 5.00%, respectively).

Conclusions Differential leukocyte analysis of the BAL fluid collected from HDM-induced asthma suggested a severe, acute invasive response, marked by increased neutrophil infiltration. In contrast, the AF extract asthma model produced a Th2-mediated chronic allergic response, as marked by increased eosinophil and lymphocyte infiltration.
Sarcopenia is associated with frailty in lung transplant candidates


Purpose of study Sarcopenia is associated with mortality in lung transplantation (LT). Sarcopenia is considered a part of frailty; this relationship has not been confirmed in lung disease.

Methods used In 367 LT candidates, we used three sarcopenia definitions: European Working Group on Sarcopenia in Older People (EWGSOP); FNIH Sarcopenia Project (FNIH); and lowest quartile of appendicular skeletal muscle mass (ASMI). ASMI was measured by bioelectrical impedance analysis. Frailty was measured by Short Physical Performance Battery (SPPB; SPPB $\leq 7$ frail) and Fried Frailty Phenotype (FFP; FFP $\geq 3$ frail). Associations between sarcopenia definitions and frailty were tested by logistic regression adjusting for age, sex, diagnosis. Non-linear associations between ASMI and frailty were tested using generalized additive models (GAMs), adjusting for age, sex.

Summary of results Sarcopenia was associated with increased risk of frailty by EWGSOP and lowest quartile definitions (table 1). By GAMs, the relationship between ASMI and frailty risk was linear (figure 1).

Conclusions Sarcopenia is associated with frailty in LT candidates by EWGSOP and distribution-based definitions. Studies are needed to optimize sarcopenia definitions and determine if reducing sarcopenia improves LT outcomes.

Abstract 148 Table 1 Sarcopenia and risk of frailty, testing three definitions of sarcopenia

<table>
<thead>
<tr>
<th>EWGSOP*</th>
<th>FNIH*</th>
<th>Lowest Quartile</th>
</tr>
</thead>
<tbody>
<tr>
<td>SPPB Odds Ratio</td>
<td>3.4 (1.7–7.0); p=0.01</td>
<td>0.9 (0.4–2.2); p=0.86</td>
</tr>
<tr>
<td>(95% CI)</td>
<td></td>
<td></td>
</tr>
<tr>
<td>FFP Odds Ratio</td>
<td>2.4 (1.2–5.0); p=0.02</td>
<td>1.3 (0.6–2.9); p=0.51</td>
</tr>
<tr>
<td>(95% CI)</td>
<td></td>
<td></td>
</tr>
</tbody>
</table>

*ASM = ASM/h2<7.0 kg/m2 in men, <5.5 kg/m2 in women
ASM:BMI<0.789 men, <0.512 women
ASM:ASM/h2<7.4 kg/m2 men, <5.9 kg/m2 women

Abstract 148 Figure 1 Association between ASMI and predicted risk of frailty using (A) SPPB and (B) FFP

Surgery II

Concurrent session

3:15 PM

Thursday, January 23, 2020

Abstract 149 Automated facial measurements provide empirical data of skin compliance

S. Rodrick, C. Lee, L. Tobing, C. Drew, S. Gupta. Loma Linda University, Loma Linda, CA

Purpose of study A recent study found that concerns about social interactions and physical appearance did not improve until three months after facial reconstructive surgery and may correlate with improved healing and the appearance of skin over time. There is no empirical data and measurements of the skin’s compliance and the degree to which facial surgical site skin deviation returns to preoperative baseline measurements. Emotrics, a machine learning technology, provides an objective tool for facial analysis. The data from this study may help providers anticipate patient concerns and tailor counseling for different time points in the preoperative and postoperative periods.

Methods used An analysis of adult patients who underwent wide excision or flap repair of facial skin cancer defects were recruited and provided consent for this study. The automated facial analysis program Emotrics automatically displays the facial image with 68 facial landmarks used to locate the perimeter of the face, interpupillary distance, and a central vertical axis of the face. These landmarks on preoperative and postoperative photographs were used to quantify facial symmetry. Measurements of facial features: brow, nasal ala, and nasal base excursion were calculated from preoperative and immediate postoperative photos and preoperative and three-month postoperative photos for analysis.

Summary of results Photographic results will be presented and demonstrate a time scale for the improvement in facial skin distortion over the healing period. Brow deviation measured at 0.84 cm immediately postoperatively compared to preoperative baseline symmetry, measured at 0.22 cm of deviation from baseline at three-months postoperatively. Decay charts for the various facial regions measured demonstrate a similar phenomenon of improvement towards baseline appearance at three months.

Conclusions Measurements of the skin’s compliance and the degree to which facial surgical site skin deviation returns to preoperative baseline measurements was captured using a machine learning technology. Further data gathered may help providers tailor counseling for different time points in the perioperative periods of facial surgery and better prepare patients for healing timeframe expectations.
base excursion were calculated from preoperative and immediate postoperative photos and preoperative and three-month postoperative photos for analysis.

Summary of results Photographic results will be presented and demonstrate a time scale for the improvement in facial skin distortion over the healing period. Brow deviation measured at 0.84 cm immediately postoperatively compared to preoperative baseline symmetry, measured at 0.22 cm of deviation from baseline at three-months postoperatively. Decay charts for the various facial regions measured demonstrate a similar phenomenon of improvement towards baseline appearance at three months.

Conclusions Measurements of the skin’s compliance and the degree to which facial surgical site skin deviation returns to preoperative baseline measurements was captured using a machine learning technology. Further data gathered may help providers tailor counseling for different time points in the perioperative periods of facial surgery and better prepare patients for healing timeframe expectations.

150 EFFICACY OF INTRA-ARTICULAR INJECTIONS WITH CORTICOSTEROIDS VERSUS KETOROLAC IN THE HIP AND KNEE

1K Jurgensmeier*, 1P Fuerst, 0D Jurgensmeier, 0D Kanz, 5J Daines. 1University of Washington School of Medicine, Seattle, WA; 2Saint Alphonsus, Boise, ID

10.1136/jim-2019-WMRC.150

Purpose of study Intra-articular steroid injections have become a mainstay of treating moderate to advanced arthritis. However, concern exists regarding associated risks and adverse effects. Non-steroidal anti-inflammatory injections, such as Ketorolac, may offer similar benefits as steroids without a similar risk profile. There is, however, a paucity of direct comparison between these two injection types. This study evaluates the effectiveness of Triamcinolone, a corticosteroid, and Ketorolac, an anti-inflammatory, in treating symptoms of moderate to advanced osteoarthritis of the hip and knee.

Methods used 110 patients (52 hips, 58 knees) with documented moderate to advanced degeneration of the hip or knee were randomized in a double-blinded study. Baseline measurements were obtained via the Visual Analogue Scale (VAS), Knee injury and Osteoarthritis Outcome Score Jr. (KOOS Jr.) or Hip dysfunction and Osteoarthritis Outcome Score Jr. (HOOS Jr.), and PROMIS Global Health Score. Patients were randomly assigned to receive Ketorolac or Triamcinolone injections given via ultrasound guidance. Follow up surveys were collected 1 week, 1 month, and 3 months after the injection date. Data was compared by two-way repeated measures ANOVA with a Bonferroni posthoc test of means comparison.

Summary of results No significant differences were noted in comparing the effectiveness of Triamcinolone or Ketorolac for the various facial regions measured demonstrate a similar time scale for the improvement in facial skin distortion over the healing period. Brow deviation measured at 0.84 cm immediately postoperatively compared to preoperative baseline symmetry, measured at 0.22 cm of deviation from baseline at three-months postoperatively. Decay charts for the various facial regions measured demonstrate a similar phenomenon of improvement towards baseline appearance at three months.

Conclusions Ketorolac is non-inferior compared to Triamcinolone in providing improved functionality and pain relief to degenerative joint disease in the hip and knee. This allows providers an additional low-cost medication to offer patients with contraindications to steroid injections. Further studies will be beneficial to assess if Ketorolac has a similar risk profile when compared to steroid injections.

151 EFFECT OF PAIN MEDICATION REGIMEN ON HOSPITAL LENGTH OF STAY AFTER CLEFT PALATE SURGERY


10.1136/jim-2019-WMRC.151

Purpose of study Optimal post-operative length of stay (LOS) for patients having cleft palate surgery is unclear. This study aims to understand how pain medication regimen affects post-operative LOS following primary cleft palate repair.

Methods used A retrospective chart review was performed on 200 consecutive patients who had cleft palate repair from June 2011 to May 2018 at BC Children’s Hospital. Data collection included: patient demographics, post-operative pain medication regimens (e.g. dose, frequency, regularity), and post-operative LOS.

Summary of results The cohort of 200 patients (108 male: 92 female) had a median age of 11.0 months (IQR 9.5 – 12.4) at the time of surgery, and a median post-operative LOS of 43.1 hours (IQR 26.1 – 50.1). Regularly dosed acetaminophen (n=77) was associated with a shorter median LOS of 28.6 hours compared to 45.5 hours for PRN dosed acetaminophen (n=122), (p=0.02). Regularly dosed ibuprofen (n=20) was also associated with a shorter median length of stay of 26.1 hours as compared to 43 hours for PRN dosed ibuprofen (n=75) and 43.9 hours for no ibuprofen (n=105), (p=0.03). Length of stay was not associated with patients receiving only acetaminophen (n=104) or both acetaminophen and ibuprofen (n=95) (p=0.26). LOS was not associated with the type of opioid analgesia prescribed: codeine (n=66), morphine (n=65), or codeine and morphine (n=58), (p=0.95).

Conclusions Patients receiving regularly dosed acetaminophen or ibuprofen have shorter LOS following cleft palate repair. Opioid analgesia is not associated with post-operative LOS.

152 DEVELOPMENT OF A MODEL TO RECRUIT T-REGULATORY CELLS TO VASCULARIZED COMPOSITE ALOOOGRAFT TISSUE

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Purpose of study Regulatory T lymphocytes (T-regs), play a key role in immune homeostasis due to their ability to suppress effector T lymphocytes. Cancer immunology studies have shown that recruitment of T-regs enables tumors to evade the immune system using C-C motif chemokine ligand 22, or CCL22, to recruit T-regs. This concept has been extrapolated...
Resuscitative Thoracotomy in the Emergency Department: Factors and Survival

Purpose of study The use of resuscitative thoracotomy in trauma patients remains a controversial topic. Previous investigations have reported that patients with blunt injuries have a higher rate of mortality than patients with penetrating injuries. The current retrospective study aimed to further explore the specific mechanisms of injury and factors that may increase survival of the patient undergoing the procedure.

Methods used All patients with thoracotomies performed at Arrowhead Regional Medical Center (ARMC) from 2011–2019 were extracted from Electronic Medical Record (EMR). Patients were excluded from the analysis if they were less than 18 years old. Parameters of interest included mortality, age, time of thoracotomy compared to time of arrival, and mechanism of injury (blunt versus penetrating). Data was extracted from nurse notes, paramedic notes, discharge summaries, and EMR.

Summary of results Among the 40 patients included in the original database, four patients were excluded due to age <18 years. As a result, 36 patients were included in the final analysis. The average age was 38.2 (SD=15.1) years, 75% (n=27) were males, 55.6% (n=20) were Hispanic, 41.7%, 69.4% (n=25) sustained penetrating trauma, and the average Injury Severity Score (ISS) was 27.6 (SD=17.2). The overall mortality was 77.8% (n=28). Blunt trauma was associated with 100% mortality (n=11 of 11), while penetrating trauma was associated with a 68% mortality (n=17 of 25), however the difference failed to reach statistical significance due to small sample size (p=0.0757). Factors such as age, gender, and ISS were not statistically significantly associated with mortality.

Conclusions The primary aim of the study, which was to analyze thoracotomy survival rates at a regional trauma center based on a number of demographic and other factors, was met. The knowledge provided by this study highlights the need for continual revisiting of thoracotomy guidelines in order to decrease utilization of the procedure based on circumstance. Further multicenter studies are needed to illuminate how mortality is impacted by the many external factors listed previously.

Impact of Intraoperative Indocyanine Green Angiography (SPY-PHI) for Patients with Facial Defects on Operative Time and Number of Procedures

Purpose of study Basal and squamous cell skin cancers present a multiplex problem because of the risks of major disfigurement and muscle or nerve injury, especially in fragile areas such as the face. This causes an increase in operative time and trips to the operating room for patients with these facial defects. However, SPY Portable Handheld Imager (SPY-PHI) is a tool that uses indocyanine green fluorescence imaging technology that could aid in reducing operation time and trips. SPY-PHI enhances the surgeon’s ability to assess tissue viability by intraoperatively evaluating blood flow and tissue perfusion; thus, providing a more efficient method.

Methods used This is a case series using intraoperative SPY-PHI angiography to evaluate patients with facial defects such as basal and squamous cell skin cancers. SPY-PHI was used to assess blood flow, examine the extent of vascular injury, and assist in flap tailoring. Angiography with SPY-PHI is performed at the beginning of the case to demarcate non-viable tissue and affected areas are marked for excision. It is subsequently used after flap elevation to ensure perfusion of transported tissue. Total operative minutes and trips to the operating room are compared to historical cases where SPY-PHI technology was not used.

Summary of results Out of 30 patients, 4 have been assessed with intraoperative SPY-PHI angiography. With SPY-PHI, the average number of trips to the OR was 2.75 and the average operation time was 577.5 minutes. For the 26 patients that did not undergo SPY-PHI, the average number of trips to the OR was 1.35 and the average operation time was 202.08 minutes.

Abstracts
Conclusions The previous hypothesis stated that SPY-PHI technology would decrease the time spent in the OR as well as trips to the OR. However, this was rejected as the results revealed that using SPY-PHI intraoperatively took more operating time and trips for patients with basal and squamous cell skin cancers. A possible reason for this outcome is that the cases which had used SPY-PHI were more complicated compared to those that had not used intraoperative angiography. Nonetheless, although SPY-PHI is highly useful in aiding surgeons to accurately assess tissue injury and viability, it does not reduce operative minutes and trips to the OR.

10.1136/jim-2019-WMRC.155

**EVALUATION OF COHORT PROGRESSING FROM FIRST TO SECOND STAGE SACRAL NEUROMODULATION AND INDICATIONS FOR UNPLANNED DEVICE REMOVAL**

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Purpose of study Sacral neuromodulation (SNM) is indicated for the treatment of overactive bladder, urinary retention, and bowel disorders, and is typically implanted in two procedures. Patient selection criteria that predict progression from the first to second Stage of implantation is not well defined. The purpose of this study is to determine whether symptoms, comorbidities, and demographics influence progression to second Stage implantation or future unplanned SNM removal or revision. This study seeks to provide additional clinical guidance for deciding between staged or direct full (single) Stage procedures.

Methods used A retrospective review was conducted in patients who underwent staged SNM at a single hospital by five different providers between 2012 and 2019. Outcome was measured as (1) progression from Stage 1 to 2, and (2) indications for unplanned SNM removal or revision. Chi Square analysis, Mann-Whitney U, and Fisher’s exact tests were used for data evaluation.

Summary of results A total of 153 patients underwent SNM therapy for symptoms of overactive bladder (n=129), urinary retention (n=42), neurogenic bladder dysfunction (n=18), fecal incontinence (n=18), and constipation (n=5). 92.2% (n=118/128) of patients progressed to Stage 2. Diagnoses of urinary retention and neurogenic bladder dysfunction were associated with lower progression to Stage 2 (p=0.034, p=0.017, respectively). Non-obese patients (BMI <30 mg/kg²) were more likely to have SNM removal or surgical revision within 4 years (17.5%) than obese (BMI ≥ 30 mg/kg²) patients (9.5%) (p=0.041). Other demographic characteristics and medical parameters were not associated with SNM outcome.

Conclusions The high progression rate to Stage 2 supports the concept that direct full Stage SNM can be performed in order to save time, costs, and potentially reduce morbidity related to an additional surgery. However, staged procedures should be discussed with patients with urinary retention and neurogenic bladder dysfunction due to their lower progression rates. BMI may also be an important selection criterion since a lower BMI is associated with a higher rate of SNM removal or revision. Future prospective studies are needed to further explore direct full Stage SNM implantation and parameters that predict SNM outcome.

**THE IMPORTANCE OF PREOPERATIVE NUTRITION IN PLASTIC SURGERY PATIENTS: A META-ANALYSIS OF KEY NUTRIENTS**

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10.1136/jim-2019-WMRC.156

**TRIAL OF A VIRTUAL REALITY TOOL TO TEACH SURGICAL TECHNIQUE FOR TIBIAL SHAFT FRACTURE INTRAMEDULLARY NAILING**

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10.1136/jim-2019-WMRC.157

Purpose of study The aim of this study was to compare the transferability of skills gained in a VR training program of a simulated intramedullary nailing (IMN) of a tibia by comparing performance of VR trained and standard guide (SG) trained novice medical students when performing a SawBones simulation of intramedullary nail fixation.
Methods used 20 first- and second-year novice medical students were recruited and randomized into VR (n=10) and SG (n=10) groups. Students with prior knowledge of the procedure were excluded. All 20 participants completed the first phase and 17 completed the second phase of the study. Participants were randomized to SG or VR training. After training, participants were then observed performing the tibial IMN procedure with SawBones and evaluated by a blinded attending surgeon using procedure-specific checklist and 5-point global assessment scale. Participants returned after 2-weeks for repeat training and evaluation.

Summary of results Aggregate global assessment scores were significantly higher for VR than SG group (17.5 vs. 7.5, p<0.001), as were scores in all individual categories. The percentage of steps completed correctly was significantly higher in the VR group compared to the SG group (63% vs. 25%, p<0.002). The average improvement between the first and second phases of the study were higher in the VR group compared to SG group across all 5-categories of the global assessment scale, and significantly higher for knowledge of instruments (50% vs. 11%, p <0.01).

Conclusions Virtual reality training was more effective than standard training in our model of simulated tibia IMN for novice medical students. Virtual reality training may be a useful and cost-effective method to reduce the initial learning curve and augment orthopaedic surgery training.

Poster session
Adolescent medicine and general pediatrics
6:00 PM
Thursday, January 23, 2020

159 HYDROCARBON ASPIRATION PNEUMONITIS
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Background Mineral oil, in the form of baby oil, is a common household product when there are young children in the home. It is often overlooked as a potential source of harmful ingestion. This case is presented to raise awareness that
everyday items in the home can cause significant injury to young children. Case This is a previously healthy 16 month old female who presented to the pediatric emergency department (PED) with fever and increased work of breathing after ingesting an unknown amount of baby oil. The night prior to arrival, parents witnessed the child drink an opened bottle of baby oil. Overnight, she had a continuous cough, and the next day noted to have increased respirations and fever. She was taken to the PED, and was found to be febrile to 101.8 F and tachycardic, but with a benign lung exam. Her heart rate soon normalized and her lungs remained clear. Labs showed leukocytosis, elevated AST, and CRP. Chest XR demonstrated scattered airspace disease bilaterally more extensive on the right. Poison control recommended 24 hour observation and antibiotics. The patient was admitted for observation and continued on IV clindamycin. She remained afebrile and hemodynamically stable on room air. She continued to improve clinically and was discharged the next day without antibiotics. Discussion Hydrocarbon ingestion primarily exerts its effects on the respiratory and central nervous systems. Most concerning and most often seen are the respiratory complications that result from aspiration because of the organic properties of hydrocarbons. Pulmonary symptoms occur within 30 minutes of ingestion and include coughing, choking, tachypnea, dyspnea, rales, and/or grunting. Aspiration into the lung parenchyma leads to chemical pneumonitis and direct injury to the lung tissue. The prognosis for children who have ingested a hydrocarbon is dependent on the amount and specific agent involved. While most children survive without any complications, some can quickly progress to respiratory failure and death. The importance of this case is to remember that simple, common household products can cause major injuries to children. As a result, pediatricians need to include these products in their anticipatory guidance about poisonings, in addition to the more routinely discussed substances.

NEONATAL DIABETES MELLITUS

5 Chen*. UCSF-Fresno, Fresno, CA

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Introduction Neonatal diabetes mellitus is characterized by persistent hyperglycemia typically within the first 6 months of life due to impaired insulin function. Etiology of neonatal diabetes is mostly monogenic and is an extremely rare condition worldwide with an estimated incidence of 1:300k to 1:500k individuals. We present a case of neonatal diabetes mellitus that was able to respond to dietary regulation without insulin therapy.

Case presentation An ex-37 6/7 wk newborn presented with recurrent episodes of hyperglycemia. Patient was first noted to have hyperglycemia on DOL 1. Glucose at birth was 159 but had quickly risen to 230. Patient was also noted to be hypotonic and having some temperature instability. Patient was transferred to the NICU at that time and was noted to have increased respirations and fever. She was taken to the PED, and was found to be febrile to 101.8 F and tachycardic, but with a benign lung exam. Her heart rate soon normalized and her lungs remained clear. Labs showed leukocytosis, elevated AST, and CRP. Chest XR demonstrated scattered airspace disease bilaterally more extensive on the right. Poison control recommended 24 hour observation and antibiotics. The patient was admitted for observation and continued on IV clindamycin. She remained afebrile and hemodynamically stable on room air. She continued to improve clinically and was discharged the next day without antibiotics. Discussion Hydrocarbon ingestion primarily exerts its effects on the respiratory and central nervous systems. Most concerning and most often seen are the respiratory complications that result from aspiration because of the organic properties of hydrocarbons. Pulmonary symptoms occur within 30 minutes of ingestion and include coughing, choking, tachypnea, dyspnea, rales, and/or grunting. Aspiration into the lung parenchyma leads to chemical pneumonitis and direct injury to the lung tissue. The prognosis for children who have ingested a hydrocarbon is dependent on the amount and specific agent involved. While most children survive without any complications, some can quickly progress to respiratory failure and death. The importance of this case is to remember that simple, common household products can cause major injuries to children. As a result, pediatricians need to include these products in their anticipatory guidance about poisonings, in addition to the more routinely discussed substances.

Management and outcome Endocrinology was consulted. Patient was found to have low/undetectable levels of insulin. Patient was suspected to have neonatal diabetes mellitus. Genetics was consulted as monogenic diabetes is the most common cause of neonatal diabetes. Patient remained in the NICU for glucose monitoring. Insulin therapy was not initiated as patient does occasionally dip to the 100’s and risks of hypoglycemia with insulin therapy outweighed risks associated with hyperglycemia. Patient feeding frequency was increased to Q2 hr to help stimulate pancreas to keep up with carbohydrate intake. Patient’s blood glucose began improving to the 130’s to 170’s. Although blood glucose remains elevated, patient was stable enough for discharge home on DOL 18. Patient scheduled to have close follow up with genetics and endocrine following discharge.

Discussion The decision to start insulin therapy always requires careful consideration as risk of hypoglycemia in neonates can lead to poor neurocognitive development in the future. This case demonstrated an interesting presentation of neonatal diabetes mellitus which responded to non-pharmacologic management and dietary regulation alone. This case will add to the understanding of neonatal DM and aid in the decision for initiation of insulin therapy in future cases.

161 ESTABLISHING GUIDELINES FOR PEDIATRIC FEVER IN A COMMUNITY HOSPITAL

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10.1136/jim-2019-WMRC.161

Purpose of study Fever is one of the most common complaints for pediatric emergency department (ED) visits. As many pediatricians know, serious bacterial infections (SBI) can be difficult to diagnose as infants may only present with fever. Multiple studies have shown that prompt identification and administration of appropriate empiric antibiotic therapy prevents significant morbidity and mortality and improves outcomes. Working in a large, level 1 trauma community hospital with an annual ED volume of 110,000 patients, often pediatric patients are alongside acutely decompensating adult patients resulting in delay of care. This study aims to establish a pathway to aid in appropriate management of these patients.

Methods used A needs assessment among physicians and nurses in the ED and department of pediatrics led to the creation of a new ‘Pediatric Fever’ pathway. The primary outcomes for analysis include (1) appropriate identification of high risk pediatric patients with a fever (2) administration of empiric antibiotics within an hour of identification of a fever in the ED and (3) time to admission from the ED. The algorithm includes transition of care from the ED to inpatient setting. The study also involves retrospective chart review of patients age 72 hours to 2 years presenting with fever, and use of the above measures to compare before and after implementation of the pathway for 3 months.

Summary of results The implementation of the Pediatric Fever Pathway in our level one trauma ED and pediatric inpatient setting will provide insight on the percentage of febrile pediatric patients that were appropriately given antibiotics within one hour of identification of fever. The data will also reflect whether the appropriate antibiotic was initiated in the ED.
Abstracts

Conclusions Many children’s hospitals have various pathways that aid healthcare providers. As a community hospital now caring for more children, it is imperative we collaborate with ED physicians to optimize care. This pathway hopes to address errors in identification and management of pediatric SBI. It promotes antimicrobial stewardship by avoiding inappropriate use of broad spectrum antibiotics to minimize bacterial resistance. This pathway also allows for education of resident doctors and a collaborative approach among emergency medicine and pediatrics.

162 THE PREPARED FIRST RESPONDER: ENGAGING YOUTH ADVOCATES AT SCHOOL, HOME AND IN THE COMMUNITY

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Purpose of study With youth empowerment at the core of this project and to engage with our surrounding community we found an opportunity to actively work with at-risk Fresno youth on their terms. We partnered with a local high school, Design Science Middle College High School, to develop interactive educational modules in educational areas that the youth found interesting, attributable to their daily living and where they felt they could genuinely advocate for change as advocates at school, home and in their community.

Methods used A PhotoVoice project was implemented as a needs assessment tool into the tenth-grade class at DSMCHS to better understand their needs. Three modules were created focusing on topics our students found important; first aid, nutrition, and youth engagement. Surveys were administered before and after the modules using Qualtrics.

Summary of results Twenty-seven percent of students reported being concerned about gang violence, drugs/alcohol, and/or poverty within their communities. "[l]ife after college" and "[h]aving a place to belong and feel safe" were their two largest stressors. Sixty-nine percent reported consumption of sodas or sugar-sweetened beverages 1–2 times a day. Seventy percent reported feeling more confident in writing a policy that can positively impact their community.

Conclusions During the educational modules, students acquired first aid skills to be able to respond to acute injuries they might encounter in everyday life learned how to read food labels to make healthier choices and prepared health presentations to improve their public speaking skills.

163 ALTERED EXPRESSION OF THIOREDOXINS, PEROXIREDOXINS, AND THIOREDOXIN REDUCTASES IN CBS-DEFICIENT HOMOCYSTINURIA IN THE PRESENCE AND ABSENCE OF HOMOCYSTEINE-LOWERING TREATMENT: POSSIBLE IMPLICATIONS FOR REDOX MEDIATED PATHOLOGY

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Purpose of study Cystathionine β-synthase-deficient homocystinuria (HCU) is a poorly understood, life-threatening, inborn error of sulfur metabolism. If left untreated, it can lead to cognitive impairment, connective tissue disturbances, and thromboembolic complications. Multiple lines of evidence from both the transgenic HO mouse model of HCU and human HCU patients indicate oxidative stress as a major pathogenic factor in this disease. Previous work has shown that impaired antioxidant defense may contribute to the generation of oxidative stress in HCU.

Methods used We investigated the hepatic expression of the antioxidant proteins thioredoxin (TRX) 1, thioredoxin reductases (TRD) 1 and 2, and peroxiredoxins (PRDX) 1,2 and 3 in the presence and absence of the homocysteine-lowering therapy betaine in the HO mouse model of HCU.

Summary of results Western blotting analysis revealed significant repression of TRDX, TRD1 and TRD2 in untreated HCU mouse livers. These effects were not ameliorated by betaine treatment. PRDX 1, 2 and 3 were all induced in HCU livers. Betaine treatment normalized PRDX 1 and 2 expression levels but induced a further induction of PRDX3.

Conclusions Our data suggests that impaired expression of thioredoxin 1 and thioredoxin reductases has the potential to contribute to oxidative stress in HCU and that the induction of multiple peroxiredoxin isoforms may constitute a redox-sensitive homeostatic response in this disease.

164 CLINICAL PATHWAY DEVELOPMENT FOR CHILDREN WITH A SOLITARY FUNCTIONING KIDNEY

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Purpose of study Congenital anomalies of the kidney, including a solitary functioning kidney (SFK), are frequent causes of childhood chronic kidney disease (CKD). Those at high risk of developing CKD likely require a different standard of long-term care. We hypothesized that long-term SFK outcomes are not uniformly favorable, extensive local practice variation exists for length of follow-up and extent of surveillance, and risk stratification may improve care efficiency.

Methods used This was a retrospective cohort study. All patients identified with a SFK from 2000–2017, due to either renal agenesis (RA) or multicystic dysplastic kidney disease (MCDK), were included. Relevant data was extracted from clinical charts. Outcomes highlighting local care patterns and
resource use included length of follow-up, and number of clinic visits and renal ultrasounds over total follow-up. To assess the utility of risk-stratification, SFK patients were stratified into high (HR) and low risk (LR) groups based on an estimated glomerular filtration rate (eGFR) <90 ml/min/1.73m² or ≥90 at last follow-up, respectively. Between-group differences were analyzed by Mann-Whitney U test.

Summary of results A total of 229 SFK patients (71 RA, 158 MCDK) were included. Median age at initial visit was 0.3 (IQR 0.6) years with a median follow-up of 4.3 (4.8) years. The mean eGFR at last follow-up was 89±28 ml/min/1.73m². After stratification based on last eGFR, 53 patients (23%) were classified as HR and 176 (77%) as LR. With regards to practice variation, median number of clinic visits over total follow-up was 4 (2) ranging from 1–47, while median number of ultrasounds was 4 (4) with a range of 0–28. When practice patterns were stratified according to risk, there were no significant differences between HR and LR groups for length of follow-up (3.9 vs 5.8, p=0.07), number of clinic visits (5 vs 6, p=0.26), or number of ultrasounds (5 vs 5, p=0.57).

Conclusions Significant practice variation exists in the management of children with a SFK. Moreover, despite differences in long-term kidney function, the care of LR and HR patients is similar, suggesting that current care patterns warrant standardization and risk-based allocation of resources.

Students responded to confidential surveys midway through and at the end of the academic year that assessed their mental health using the Brief Resilience Scale (BRS), perceptions of the impact of the days, and suggested improvements. The results were tabulated.

Summary of results Of the class of 160 students, 80% filled out the final survey. Of the responders, 62% experienced a direct positive effect on their mental health from the program, 68% learned new skills, and 61% felt less alone.

There were no significant differences in the BRS scores of the aggregated students in December 2018 (3.00, n=147) or June 2019 (2.99, n=133), nor were there differences in score between gender: female (2.94, n=57) and male (3.02, n=74). These scores indicate that students at the end of their third-year have borderline low to normal resilience.

Conclusions Students reacted positively to the addition of Resiliency Days and felt that they had a beneficial impact on their mental health. They gained tools that may help them face future challenges and felt less alone. Impact on BRS scores was not seen likely because of the many factors that contribute to one’s resilience.
Cultivating mindfulness disposition may be beneficial for improving physiological profiles among smokers in withdrawal. This finding has value for future studies that incorporate mindfulness as a behavioral intervention for smokers attempting to improve their stress response and reduce their urge to smoke. Limitations include a small sample size; we aim to retest our analyses in a larger sample.

Abstract 167

EVALUATION OF DYNAMIC MONITORS FOR THE PREDICTION OF VOLUME RESPONSIVENESS IN PATIENTS WITH AND WITHOUT DIASTOLIC DYSFUNCTION

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Purpose of study Patients with diastolic dysfunction (DD) comprise more than 28% of patients aged 60 and older. Left ventricular DD is known to predict adverse outcomes such as major adverse cardiac events, and in-hospital mortality. Intraoperative optimization of cardiac function, in particular that of intravascular volume, can decrease incidence of complications. Current gold standard of measurement of cardiac function in response to intravenous fluids in patients under general anesthesia is the change in arterial waveform (PPV – Pulse Pressure Variation, SVV – Stroke Volume Variation) associated with respiration. The SVV values predictive of fluid responsiveness have been characterized in a general patient population. Thresholds haven’t been specifically validated for patients with DD. Identification and management of these patients can improve perioperative outcomes. Our research aims to evaluate and characterize the potential impact of DD in response to each fluid bolus was automatically recorded by the monitor, and comparisons were made between patients with and without DD.

Methods used Eligible participants were invited to complete the anonymous survey via the AD Facebook site with approximately 400 members. The survey instrument contained 21 questions covering demographics, history of injuries and additional questions specific to AD. Inclusion criteria: self-identified Dunkers. Exclusion criteria: age <18 years.

Results To date, 35 patients have been recruited. ROC curve analysis was produced to compare fluid responsiveness in those with or without DD.

Conclusions The total sample size is currently too small for conclusions. Additional data is necessary to fully characterize the relationships.
IMPACT OF CAFFEINE ON SKIN CANCER PREVENTION: A META-ANALYSIS

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Purpose of study Skin cancer has an annual incidence of 5.5 million in the U.S., more than all other cancers combined. Epidemiological studies show that caffeinated beverage intake is associated with decreased risks of several skin cancer types: basal cell carcinoma (BCC), squamous cell carcinoma (SCC), and melanoma. Due to the high incidence of skin cancer and the popularity of caffeinated drinks, caffeine may have an appreciable impact on skin cancer incidence. We performed a systematic review and meta-analysis to estimate the annual reduction in skin cancer incidence and treatment cost by caffeinated beverage intake.

Methods used We searched PubMed, Embase, and Scopus for observational studies that assessed the relative risk of skin cancer with coffee or tea intake up to June 30, 2019. High-quality studies were selected by the Newcastle-Ottawa Quality Assessment Scale for further analysis. Summary relative risk (SRR) was determined by the DerSimonian and Laird random effects model.

Summary of results A database search yielded 22 high-quality studies covering 47,593 cases of skin cancer, mostly in the U.S. and Europe. For all skin cancer types combined, caffeinated coffee had the largest effect with SRR (95% confidence interval) of 0.78 (0.68–0.87). Tea had less effect (0.85 (0.77–0.93)), and decaffeinated coffee had no effect (0.98 (0.91–1.06)). Combining coffee and tea intake, SRRs were 0.88 (0.81–0.95) for BCC, 0.81 (0.69–0.93) for melanoma, 0.77 (0.59–0.94) for SCC, and 0.77 (0.59–0.95) for unspecified nonmelanoma skin cancer (BCC or SCC). These effects were dose-dependent. Based on SRR, dose dependency, and proportion of coffee drinkers, we estimate that current coffee consumption leads to 196,344 skin cancers prevented and 289 million dollars saved in treatment cost annually in the U.S.

Conclusions Current caffeinated beverage intake significantly lowers the burden of skin cancer in incidence and treatment cost. This meta-analysis has limitations including varying caffeine content in coffee consumed. Future studies should clarify the ideal timing of caffeine intake relative to sunlight exposure.

A SURVEY OF PRIMARY CARE PHYSICIANS ON HYPERHIDROSIS TRAINING AND EXPERIENCE

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Purpose of study Data on hyperhidrosis education among physicians is lacking. Hyperhidrosis is defined as excessive sweating beyond the body’s needs for temperature regulation. The cause of primary focal hyperhidrosis is not well understood. Palmar and axillary primary focal hyperhidrosis are ranked as having the highest impact on quality of life when compared to 40 other dermatologic conditions. The aim of this project was to collect information on the nature and quality of training and knowledge related to hyperhidrosis within family medicine and internal medicine residency programs. In a recent study approximately 70% of hyperhidrosis patients reported seeing a primary care physician about their condition. This statistic highlights the need for adequate training of primary care physicians to manage patients with hyperhidrosis.

Methods used An 8-question survey was administered to Family Medicine (FM) and Internal Medicine (IM) residents and attending physicians within the Samaritan Health Services FM and IM programs via paper copy and Survey Monkey. The survey was sent to 32 residents and 47 attending physicians. 32 residents and 13 attending physicians responded to the survey.

Summary of results We recorded a 57% response rate. 92% of attending physicians and 53% of residents indicated that they had encountered a patient with either generalized or focal hyperhidrosis. Approximately 35% of attendings and 75% of residents reported feeling less than reasonably comfortable with diagnosing and managing a patient with hyperhidrosis. Only 7% of all respondents indicated that they were familiar with iontophoresis as a treatment option. It also appears that physicians may tend to underestimate the effect that hyperhidrosis has on an average patient’s quality of life. 89% of all respondents reported having no specific training on hyperhidrosis diagnosis or management.

Conclusions There appears to be a need for increased awareness and training for primary care physicians on understanding and managing patients with hyperhidrosis. Further work is required in expanding the survey to improve the generalizability of our results.
to our analysis based on this sample size, drive time but not distance in miles was significant, no insurance vs. commercial and public was trending towards significance, and Hispanic ethnicity was not significant for lost to specialist follow up.

**Conclusions** The key findings demonstrate that neither distance nor insurance type were significantly associated with lost to follow up in ACHD at specialty treatment sites. For future, we would like to determine if our findings are consistent in other states. With this information, we can improve our understanding of the ACHD population and work to develop a system to reduce loss of care in the ACHD population.

**Poster session**

**Community health and global health**

6:00 PM

Thursday, January 23, 2020

**172 PILOT STUDY: USING CENTERING DIABETES FOR ADULT PATIENTS WITH TYPE II DIABETES AT A FREE CLINIC**

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10.1136/jim-2019-WMRC.172

**Purpose of study** Diabetes is a multisystem disease, disproportionately affecting minorities, that can lead to complications such as heart disease, stroke, kidney failure, lower limb amputation, and adult-onset blindness if not adequately controlled using the standard of care guidelines. The objective was to examine the use of diabetes group appointments using a Centering model in a free clinic setting. Patients attending Yakima Union Gospel Mission are often inadequately treated for their diabetes care model and enabled a small group of patients to meet diabetes care guidelines. Challenges were met due to patient availability and difficulty recruiting patients. Patients were satisfied with the group appointments and agreed to continue attending centering appointments based on survey data. While statistically significant differences to blood pressure and HbA1c cannot be drawn from the data, community acceptance of the centering model has opened doors to further studies.

**Summary of results** Three patients consistently participated in Centering Diabetes appointments: a 44 y.o female with a 10 pack-year history of smoking; 49 y.o. male with difficulty maintaining weight due to insulin use; and a 47 y.o female with significant work and spouse stress. Pre- to post-Centering HbA1c were 12.1 to 11.2, 12.3 to 11.9 and 9.6 to 8.4, while pre- and post-Centering blood pressures were 154/96 to 120/70, 150/100 to 145/100, and 143/80 to 130/76 respectively.

**Conclusions** The curriculum topics addressed the standard of care model and enabled a small group of patients to meet diabetes care guidelines. Challenges were met due to patient availability and difficulty recruiting patients. Patients were satisfied with the group appointments and agreed to continue attending centering appointments based on survey data. While statistically significant differences to blood pressure and HbA1c cannot be drawn from the data, community acceptance of the centering model has opened doors to further studies.
on the validity of task-sharing in surgery with non-clinician surgeons and the scaling-up of innovations addressed in the literature will help realize timely access to Bellwether procedures.

174 MITIGATING ALCOHOL ABUSE IN THE BOOMTOWN ECONOMY OF DOUGLAS, WYOMING
A Blaine*. University of Washington School of Medicine, Seattle, WA
10.1136/jim-2019-WMRC.174

Purpose of study As a community that revolves around the energy industry, Douglas, WY is home to transient employees. The Douglas community maintains an above average excessive drinking rate and a high rate of alcohol impaired driving deaths. Community conversations and further evaluation of community assets informed that Douglas needs support services to reduce alcohol consumption among transient energy industry employees.

Methods used County health data proved a significant increase in excess drinking as well as an increase in the rate of alcohol impaired driving deaths in Douglas. Solutions for Life (SFL) a state subsidized mental health clinic and local physicians were interviewed regarding the local public health concern. An evidence based literature review outlined primary and secondary factors influencing alcohol abuse. Interviews with various stakeholders in the community and clinical observations provided further insight about energy industry employees perception of how much alcohol consumption is normal. Community conversations also considered the social environment and living conditions of transient employees.

Summary of results Evidence based literature identified alcohol abuse interventions that focused on rural communities, transient employees of the energy industry, cultural alcohol misconceptions, social services, and collaborations with energy corporations. The director of SFL confirmed an excessive number court ordered patients in treatment were from the energy industry. Physicians and SFL professionals confirmed that energy employees were living in hotels and temporary housing with poor social networks. Considering the demographic of alcohol abuse reported by the community, evidence based literature determined that a targeted community health campaign towards energy industry employees in collaboration with energy corporations would be the most effective intervention to reduce alcohol consumption in rural communities.

Conclusions Support through a strength-based approach from community assets such as SFL provided an understanding of the social background in Douglas. Next steps would include the development of collaboration between SFL and energy corporations to target energy industry employees. The collaboration would facilitate a community health campaign with the intention to decrease alcohol abuse in Douglas.

175 MULTIDISCIPLINARY SUPPORT FOR BUPRENORPHINE-NALOXONE TREATMENT OF OPIOID USE DISORDER
A Hall*. University of Washington School of Medicine, Seattle, WA
10.1136/jim-2019-WMRC.175

Purpose of study Opioid use disorder (OUD) is a significant concern within the population served by the Confluence Health system in Wenatchee, Washington with an average of 17.6 associated deaths annually. This project utilized an asset-based approach to examine fundamental resources, obstacles, and potential improvements in the prescription of buprenorphine-naloxone for OUD in the community.

Methods used Discussion with a buprenorphine-naloxone prescriber outlined pharmacological therapy for OUD available in Wenatchee. OUD patients receiving buprenorphine-naloxone treatment were interviewed regarding craving control, use of addiction sponsors, therapy and support groups. The clinical director of an addiction treatment center detailed outpatient services for OUD and an addiction group therapy session was attended. Meeting with a team responsible for a buprenorphine prescriber incentive program validated administrative support for OUD.

Summary of results Effective pharmacotherapy for OUD was determined to focus on harm-reduction and compliance. Observation of group therapy demonstrated limitations of medical management without psychological and social support. Dialogue with involved professionals allowed for distribution of other providers’ viewpoints. A poster integrating these findings was shared with buprenorphine prescribers. In assessing next steps, literature review revealed buprenorphine implants to be an effective alternative route of administration. Availability of buprenorphine implants in Wenatchee could improve medical management of OUD in patients with accessibility and diversion issues. Collaboration between mental health professionals and buprenorphine prescribers to determine the best method of administration would be necessary. The existing professional relationships in Wenatchee would foster this collaboration and improve patient-centered care.

Conclusions This evaluation integrates expertise of physicians and addiction professionals while providing a testament of buprenorphine implantation as an attainable advancement in OUD treatment given the present infrastructure in Wenatchee.

176 UNINTENTIONAL INGESTION OF MARIJUANA IN ADULTS: A CASE SERIES
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10.1136/jim-2019-WMRC.176

Purpose of study Marijuana use has become increasingly popular as more US states legalize the substance. A greater number of people are admitted to the ED due to marijuana toxicity. Our study reviews a case of adults who were unintentionally exposed. This is novel because most of the current literature focuses on marijuana poisoning in pediatric patients.

Methods used We conducted a retrospective analysis of twelve subjects. Subjects were evaluated in the Emergency Department and referred to Medical Toxicology Service after ingesting marijuana from food at a family event. Six of the subjects consented to be interviewed about their experiences which
were qualitatively analyzed and grouped into common themes. The study was approved by the Institutional Review Board.

Summary of results Three subjects required observation due to persistent symptoms. Eleven subjects tested positive for tetrahydrocannabinol (THC) via urine drug immunoassay. Two subjects tested positive for ethanol in their blood. Common symptoms experienced included confusion, difficulty speaking, weakness, nausea, tremors, and hallucinations. All subjects reported sleepiness. Subjects also reported multiple emotions, including anger, confusion, disbelief, and helplessness. Three of the interviewed subjects reported a negative impact on work.

Conclusions This case series illuminates adverse effects after unintentional marijuana exposure in adults. More incidents of unintentional marijuana intoxications are expected as legal availability increases. Public education of marijuana’s effects should be widely disseminated to improve individual awareness and encourage policy implementation and regulation. This may minimize future unintentional intoxication cases.

SUPPORTING CALIFORNIA’S MEDICAL THERAPY PROGRAM SERVING CHILDREN WITH DISABILITIES THROUGH TELEMEDICINE

UC Davis Health, Sacramento, CA
10.1136/jim-2019-WMRC.177

Purpose of study California’s Medical Therapy Program (MTP) provides physical therapy (PT), occupational therapy (OT), and medical oversight to children with qualifying disabilities, generally due to neuromuscular and musculoskeletal conditions. The MTP serves nearly 23,000 children at school-based clinics called Medical Therapy Units (MTUs). While every county is required to have a MTU, not every county has a MTU, creating difficulties for children in those counties to access care. Therefore, the School-Based Tele-Physiatry Assistance for Rehabilitative and Therapeutic Services (STARS) program was developed at UC Davis Health. STARS is an innovative program that uses telemedicine to provide medical direction at the MTUs.

Methods used A needs assessment of the MTP was performed in all 58 counties through an e-mail questionnaire sent to the MTP’s Supervising Therapists. Data were collected on the counties’ resources, such as the number of MTUs. Telemedicine services began in San Joaquin and Butte Counties at the beginning of 2019. To evaluate its potential to address the shortcomings of the MTP, parent, physician and therapist satisfaction surveys were administered at the end of each monthly clinic.

Summary of results Of California’s 58 counties, 14 do not have MTUs, forcing approximately 105 children to travel to other counties for MTP services. In counties with at least one MTU, approximately 690 children travel to receive PT and OT services because their county does not have these services. The average roundtrip driving distance for physicians traveling to provide care at MTUs, was 175 miles. Furthermore, only 45% of counties have access to a pediatric physiatrist, the specialist who provides medical direction for this patient population. Preliminary satisfaction data were collected from 64 patient-therapist-physician triads. Parents (84%), therapists (91%), and physicians (100%) reported that the quality of care provided by telemedicine was equal to in-person care.

Conclusions California’s MTP is maldistributed, causing many patients and physicians to travel sometimes long distances for care. The results of the preliminary satisfaction analysis support the STARS program as a means of increasing ease of access to care and reducing burdens on families and providers.

3D IMAGE GENERATION FOR TRANSLATION OF TEXT-BASED MEDICAL QUESTIONNAIRES

B Stothers*, A Macnab. University of British Columbia, Vancouver, BC, Canada
10.1136/jim-2019-WMRC.178

Purpose of study Globally those with limited language comprehension or literacy, face problems completing validated written questionnaires related to their health/treatment, hence the importance of developing picture-based scores e.g. facial images to quantify pain. Since health care providers worldwide need software to generate images for such scales, and must choose from several options we reviewed programs suitable for first time users.

Methods used We evaluated 3 software programs (Poser version 11.0, Blender version 2.8 and Cheetah3D version 7.0) in order to select one to use to generate lifelike figures for a picture-based version of the 10-item assessment of limitations of activities (LoA) section of the SF-36 questionnaire, a validated written patient self-assessment tool. All 3 provide development tools and templates for creating figures with defined postures, facial expressions and clothing suitable for use in medical symptom scoring scales and picture-based questionnaires.

Summary of results Poser software lets users who are beginners produce figures that are more realistic and lifelike than the other programs. In particular, the ‘Smooth Translation of Joints’ feature allows more accurate rendering of human anatomy. Images can also be animated and exported in BMP, JPEG and TIFF formats. In comparison, Blender is a free high-end open source product but has a steep learning curve and while Cheetah3D is versatile, aimed at amateur artists and easy to use it is specifically written in Cocoa for MacOS users. Limitations of Poser include its cost; the inability to import artwork from other programs; and some illustration tools found in comparable software are lacking, but it does have pencils, markers, airbrushes and shading tools.

Conclusions Choices exist for those creating images for picture-based scales and questionnaires. Poser software enabled
DOES SNACKING HELP CURB CHILDHOOD OBESITY?

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10.1136/jim-2019-WMRC.179

**Purpose of study** According to the California Department of Education, students who are in Grade levels 5, 7, and 9, are near 48% overweight or obese. This is more than double what the national prevalence according to the Center for Disease Control. Unhealthy weight for this study is defined as overweight >85% BMI and obesity >95%BMI and is a serious problem that can lead to poor health, starting earlier in children. This study aims to assess a possible correlation between boredom and snacking to overweight and obesity using Body Mass Index (BMI).

**Methods used** Children between the ages of 9–15 years were referred by a physician from a local federally qualified health center in San Bernardino to ‘Operation Fit’, a weeklong day camp, based on their unhealthy weight (BMI>85th percentile). In this program, 113 children participated in a group that was taught with interactive nutrition and physical activity lessons. Parents participated in a survey for lifestyle practices, including a question about boredom and snacking, with a parent education session at the end of the camp. These answers were then compared to their child’s BMI using logistic regression models.

**Summary of results** In response to the statement ‘If I am bored, I will snack more’ there was a statistical significance for those that responded they do not snack.

**Conclusions** Those that agree with snacking when bored are 0.401 times more likely to have children that are overweight >85% BMI and obesity >95%BMI and is a serious problem that can lead to poor health, starting earlier in children. The study concludes that healthy snacking habits can be used to help lower overweight and obese children, helping with reducing the additional risks with being overweight.

**Poster session**

Endocrinology and metabolism

6:00 PM

Thursday, January 23, 2020

A UNIQUE CASE OF UNDIAGNOSED GRAVES’ DISEASE PRESENTING AS ACUTE TRANSIENT PARALYSIS

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10.1136/jim-2019-WMRC.180

**Purpose of study** Thyrotoxic Periodic Paralysis (TPP) is an unusual complication of hyperthyroidism, and is most commonly seen in Asian males. We are describing a case of a young Hispanic male presenting with acute onset lower extremity paralysis after strenuous physical activity who was found to have severe hypokalemia secondary to underlying undiagnosed Graves’ disease.

**Methods used** Retrospective case report.

**Summary of results** A 33-year-old Hispanic male with no significant past medical history presented with a two-day history of sudden onset bilateral lower extremity weakness leading to inability to ambulate. Symptoms started after patient performed strenuous activity in the setting of a sedentary lifestyle. Upon presentation to the ED, patient’s vitals were within normal limits and physical exam showed predominantly proximal bilateral lower extremity weakness, hyperreflexia in bilateral knees and 2 beats of clonus in bilateral ankles, action and postural tremors in bilateral hands. Laboratory assessment revealed severe hypokalemia, hypomagnesemia, mild elevation of creatinine phosphokinase, mild transaminitis, suppressed thyroid stimulating hormone and elevated free thyroxine. Urine potassium/creatinine ratio was less than 1 indicating transcellular shift as the cause of severe symptomatic hypokalemia. ECG 12-lead showed diffused flattened T-waves. After normalization of potassium and magnesium, the paralysis and ECG abnormalities resolved within the first day of hospitalization. Treatment for hyperthyroidism with methimazole and propranolol was started during hospitalization and patient was discharged with outpatient follow up in endocrinology clinic.

**Conclusions** This is a rare and interesting case of Thyrotoxic Periodic Paralysis in a young Hispanic male as the first manifestation of undiagnosed Graves’ disease. Thyrotoxic Periodic Paralysis often goes unrecognized at first attack due to low prevalence of disease and because patients usually exhibit mild symptoms of hyperthyroidism. TPP should always be considered in young patients presenting with acute onset muscle weakness and hypokalemia.
A CASE OF ACUTE PANCREATITIS WITH REFRACTORY ELECTROLYTE IMBALANCE

J Bhandolah*, J Bhaiya, R Janiwala, R Gupta, S Allday, C Anderson. Kern Medical, Bakersfield, CA
10.1136/jim-2019-WMRC.181

Purpose of study Pancreatitis secondary hypertriglyceridemia occurs in 1–14% of all acute pancreatitis cases. There have been no large randomized trials that have evaluated the effect of triglyceride lowering for primary prevention in patients with mild-to-moderate hypertriglyceridemia. It is important to recognize in order to provide appropriate therapy.

Methods used Retrospective case report.

Summary of results A 36-year-old male presented to the emergency department with epigastric pain with associated N/V x 1 week. Patient had a hx of alcohol abuse of 16 12-ounce beers/day x 1 yr. On physical examination, patient was in significant distress due to epigastric pain with guarding on abdominal palpation. Due to his hx and symptoms, a diagnosis of acute pancreatitis was suspected and patient was admitted for management of abdominal pain and intractable N/V. Initial lab testing did not result as patient's blood sample was found to be grossly lipemic. This provoked a lipid panel to be drawn. Lipid panel was remarkable for triglycerides of 5700 mg/dl and total cholesterol of 658 mg/dl. Of all familial hypercholesterolemia’s, Type V dyslipidemia typically presents in adulthood with high risk of acute pancreatitis. Treatment was initiated with IV Insulin and dextrose, similar to the management of diabetic ketoacidosis, although our patient was not diabetic. Blood glucose checks were done Q1 hr to adjust insulin infusion accordingly. Triglyceride levels were checked Q12 hr. Insulin enhances the activity of VLDL and Hormone-Sensitive Lipase to promote the breakdown of FFA and triglycerides in the circulation. Gemfibrozil was added to increase HDL production thus further removing triglycerides from the circulation. Outpatient management includes atorvastatin, fenofibrate, niacin, and fish oil. Lifestyle components play a major role such as alcohol cessation, avoiding complex carbohydrates, initiating an aerobic exercise plan, and increased consumption of fish containing high amounts of omega-3 fatty acids.

Conclusions There is limited data regarding which patients with hypertriglyceridemia require treatment and which therapies provide the best outcomes. This patient with pancreatitis secondary to an elevated triglyceride level of 5700 mg/dl was started on an insulin drip therapy for 3 days which lowered the level to 173 mg/dl.

Poster session
Gastroenterology
6:00 PM
Thursday, January 23, 2020

ACCUTURAL AFFECTS REPRODUCTIVE HEALTH KNOWLEDGE AND PERCEPTION OF ARMENIAN AMERICANS

A Boodaghian*, A Hovssepian, AL Nelson, F Dong. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Pomona, CA
10.1136/jim-2019-WMRC.182

Purpose of study The purpose of this study was to assess the perceptions about and knowledge of birth control amongst first and second generation Armenian men and women. A secondary objective was to investigate a possible association between education level and knowledge about birth control between the two groups.

Methods used An anonymous online survey was administered to men and women who were first and second generation Armenians from ages 18–60. The questions included information about the demographics, self-assessment (perceptions) of their knowledge about birth control and test questions designed to objectively assess their factual knowledge about contraceptive efficacy and safety. The survey was conducted in private at an Armenian community center in Burbank, CA, from 6/10/19 to 7/26/19. Chi-square tests were conducted to assess the associations between knowledge (yes vs. no) and demographic variables collected.

Summary of results Of the 147 individuals we approached, 135 participated in our study (response rate=92.8%). Overall, 61.6% were female, 33.3% were age 18–25 and 37.7% were age 26–35; 41.6% had been born in the U.S.

We analyzed the relationship between prior education and perceived and tested knowledge scores by birthplace and age of immigration. Perception of knowledge was higher than actual knowledge (p=0.04). Of those who believed they had been educated on birth control, only 33.3% answered the knowledge questions correctly, while those who believed they were not educated on the topic, 29.4% answered correctly. Higher education level correlated with higher actual knowledge (p=0.01), although those with less education perceived more knowledge on the topic.

Conclusions Overall, only a minority of respondents correctly answered questions about contraceptive efficacy. Correct test scores (knowledge) were statistically significantly associated with both higher education level and birthplace. However, perceptions of contraceptive use had no significant relationship to age, education level or acculturation, either in unmarried or married couples.
Abstract 184 Table 1

<table>
<thead>
<tr>
<th>CPT Code</th>
<th>Procedures</th>
<th>Total % Change 2000-2019 Adjusted</th>
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<tbody>
<tr>
<td>17003</td>
<td>Destruction of 2-14 skin growths</td>
<td>-94.43</td>
</tr>
</tbody>
</table>
| 11721    | Removal of tissue from 6 or more finger or toe nails | -42.25%
| 17000    | Destruction of skin growths | 19.66%
| 17110    | Destruction of up to 14 skin growths | -43.96%
| 11720    | Removal of tissue from 1 to 3 finger or toe nails | -42.55%
| 11042    | Removal of skin and tissue from 20 sq cm or less | -25.13%
| 11119    | Trimming of fingernails or toenails | -20.04%
| 11105    | Removal of single thickened skin growth | -35.65%
| 17004    | Destruction of 15 or more skin growths | -52.94%
| 11730    | Separation of nail plate from nail bed | -34.12%
| 11057    | Removal of more than 4 thickened skin growths | -18.00%
| 11043    | Removal of skin and/or muscle from 20 sq cm or less | -28.59%
| 13132    | Repair of wound (2.5 to 7 cm circumference) of forehead, cheek, chin, neck, ear, shoulder, arm, hand, and/or leg | -32.17%
| 17262    | Destruction of malignant growths (1 to 2.5 cm circumference) of trunk, arm, or leg | -10.60%
| 12632    | Repair of wound (2.5 to 7 cm circumference) of scalp, ear, shoulder, arm, and/or leg | -3.46%
| 13162    | Removal of malignant growths (1 to 2.5 cm circumference) of trunk, arm, or leg | -3.65%
| 13121    | Repair of wound (1.1 to 2 cm circumference) of scalp, arm, and/or leg | -15.91%
| 11900    | Injection of up to 1 skin growths | -54.62%
| 11770    | Removal of nail | -31.77%
| 13110    | Shaving of 0.5 centimeters to 1.0 centimeters of skin growths of the trunk, arm, or legs | -18.40%
| Average  |                         | -21.44|

Methods used The most commonly utilized dermatology procedures were found by searching for integumentary procedures in the CPY 2017 PU file data. National reimbursement averages were then calculated and analyzed from 2000 and 2019 using The Physician Fee Schedule Look-Up Tool. Using the consumer price index, the averages were adjusted for inflation.

Summary of results Medicare reimbursement for included procedures from 2000 to 2019 decreased substantially (-21.44% or 1.07%/year). The largest decreases were found in ‘Destruction of 2-14 skin growths’ (-79.43%), ‘Destruction of 15 or more skin growths’ (-52.94%), and ‘Removal of tissue from 1 to 3 finger or toe nails’ (-42.55%).

Conclusions According to our data, Medicare reimbursement in dermatology has consistently declined when adjusted for inflation. As reimbursement continually decreases, Medicare patients may experience a decrease in accessibility to dermatologists and a reduced quality of care. A comprehension of these trends is essential as continued progress is made to achieve agreeable Medicare reimbursement in dermatology.

DECREASING MEDICARE REIMBURSEMENT FOR DERMATOLOGICAL PROCEDURES 2000–2019

1A Dorius*, 1M Kropelnicki, 1JR Pollock, 2IM Haglin, 2G Hanson, 3S Ochoa, 3AR Mangold. 1B Brigham Young University, Provo, UT, 2Mayo Clinic Alix School of Medicine, Scottsdale, AZ, 3Mayo Clinic, Phoenix, AZ

Purpose of study With our aging population, patients covered by Medicare will continue to grow. For this reason, understanding Medicare reimbursement within dermatology will be increasingly important. In this study, we examine reimbursement trends for common Dermatology procedures from 2000 to 2019.

LEARNING WITH LITERATURE: HOW ARTS AND HUMANITIES CAN BE USED IN MEDICAL EDUCATION

H Drake*, C Courneya. University of British Columbia, Vancouver, BC, Canada

Purpose of study Empathy is the ability to understand the feelings of another and can be used by a physician to understand a patient’s values, feelings and perspective of their experience. This is an essential component of the doctor-patient relationship. Research demonstrates that students who engage in humanities develop higher levels of empathy and emotional intelligence; however, arts can be a challenge to incorporate in to traditional curricula of medical education.

Abstracts
Abstract 185

My Glory was I had Such Friends

Methods used A student pilot project was developed to explore empathy through reading patients’ memoirs and creating reflections. Memoirs were selected based on two criteria: that they were written from the patients’ voice and secondly, that a variety of illnesses were captured. The student then depicted the patient’s story as a visual art piece, including themes from the book and representation of their illness. This was followed by a written reflection to combine, compare and contrast all of the lessons that were learned through the readings.

Summary of results Through analyzing what the authors shared, a better understanding of what patients’ experience was developed. This led to the student having more confidence with patients, integrating tools learned from the authors, and understanding patient experiences.

Conclusions This model could be adapted to a larger number of students, each reflecting in their own artistic way on the same text. A group experience provides different perspectives of the book and results in a deeper analysis.

Abstract 186

POCKETOPHTHO: DEVELOPMENT OF A SMARTPHONE-BASED EDUCATIONAL TOOL FOR LEARNING OPHTHALMIC KNOWLEDGE

N Fong*, C Bacon, Lee-Coomes, G Yiu. University of California Davis, Sacramento, CA; California Northstate University College of Medicine, Elk Grove, CA

Purpose of study To create a high-quality smartphone app for educating ophthalmic knowledge developed under supervision of U.S. board-certified ophthalmologists.

Methods used Ophthalmic content was curated from a team of board-certified ophthalmologists at the UC Davis Eye Center. De-identified ophthalmic images were obtained from these ophthalmologists and copyright-free online sources. Educational content and images were reviewed, verified, and updated by at least two independent ophthalmologists. The app was built in XCode, an integrated development environment for building apps compatible with Mac OS-X and iOS Apple operating systems, using the programming language Swift in collaboration with students from the UC Davis Computer Science Department.

Summary of results The alpha prototype was built and reviewed with a panel of ophthalmic trainees including residents and fellows for functionality and usability to improve the prototype. The app name was selected based on survey responses. Survey questions include ‘Which app evokes reliability,’ ‘Which app sounds the most user-friendly,’ and ‘Which app sounds most reputable.’ Additional educational content has been included for development of a beta prototype.

Conclusions PocketOphtho is a high-quality smartphone app for teaching ophthalmic knowledge developed in conjunction with U.S. board-certified ophthalmologists, which can be useful for residents and fellows in ophthalmology training or in preparation for the Ophthalmic Knowledge Assessment Program (OKAP) or Written Qualifying Exam (WQE).

Abstract 187

TRANSITIONING TO ADULTHOOD WITH HYDROCEPHALUS: A PATIENT’S PERSPECTIVE

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Purpose of study Hydrocephalus is a chronic neurological condition that affects around 6 in 10 000 live births and is one of the most common indications for pediatric brain surgery. The condition is fatal if left untreated, however the introduction of new operative procedures such as shunt placements has led to most pediatric patients surviving and transitioning to adulthood. However, unlike other chronic conditions such as cystic fibrosis, congenital heart disease, type 1 diabetes, etc., the transition of adolescents with hydrocephalus from pediatric to adult care is often fragmented and disjointed given the lack of attention and research in establishing appropriate guidelines and models of transfer. This is particularly concerning not only due to the prevalence of hydrocephalus among the pediatric population, but the significant increase in morbidity and mortality associated post-transition with poorly handled transfers.

Methods used The study consisted of two phases that used a series of interview and survey questions to collect qualitative and quantitative data to identify factors that challenge young adults with hydrocephalus who are transitioning/transitioned from B.C Children’s Hospital in Vancouver into adult care at Vancouver General Hospital.

Summary of results Emerging themes from the data highlighted the difficulty patients and their family members have in forging strong and familiar relationships that have been formed over the years at B.C Children’s Hospital, adapting to the new cultural environment in an adult clinic and becoming self-reliant.

Conclusions Understanding the expectations, concerns and overall input of patients is one of the many important steps that must be taken in order to build a foundation for a transition model of care that can carefully attend to the needs of patients with hydrocephalus.
ORAL SELF-EMULSIFYING DRUG DELIVERY SYSTEM FOR DETECTION OF BETA AMYLOID PLAQUES IN THE RETINA OF AD MICE

N Hashem*, AC Bakoh, S Fuchs, M Issar. Western University of Health Sciences College of Osteopathic Medicine of the Pacific, Ontario, CA

Purpose of study Alzheimer’s disease (AD) is a neurodegenerative disease characterized by cognitive deterioration and is characterized by the accumulation of amyloid plaques found in neuronal tissue. Aβ plaques can form in retinal tissue, allowing for early detection of AD. The objective of this project was to develop a self-emulsifying drug delivery system (SEDDS) for oral administration to enhance systemic absorption of curcumin via lymphatic route. Higher blood levels of curcumin equilibrate within the eye allowing absorption to Aβ plaques in retinal tissue. The intrinsic fluorescence of curcuminoids bound to Aβ plaques would be quantified by imaging.

Methods used SEDDS were developed using medium-length chain triglycerides (MCT), polyoxyl castor oil (emulsifying agent), and Transcutol to solubilize curcumin. These formulations were observed for stability, diluted, and incubated at 37°C for 24 hrs. Before testing in vitro, formulations were selected based on particle size and triglyceride ratio and the dialysis membrane method was used to test drug release in vitro. SEDDS were diluted and 0.2 ml were added into a dialysis bag which was immersed into 19 ml of citrate buffer containing 0.5% SDS. Samples were withdrawn at different time intervals until 48 hours and were analyzed for curcumin released into the medium by HPLC.

Summary of results Addition of Oleic acid (OA) to MCT containing formulation dramatically decreased the release of curcumin from the micelles by 60%. Absolute values for drug release at 37°C for formulation #S407/10 (with OA) and #S407/9 (without OA) were 18.7% and 84.1% respectively at 48 hour. Whereas, addition of poloxamer 124 (another emulsifying agent) to polyoxyl castor oil containing formulation (#S507/8) increased drug release by approximately 80%.

Conclusions A decrease in drug release for formulation #S497/10 could be due to greater lipophilicity of oleic acid, which is a long chain fatty acid. On the contrary, incorporation of a hydrophilic surfactant poloxamer 124 to polyoxyl castor oil possibly allowed greater release due to higher dissolution medium penetration within the micelles. Oral delivery of SEDDS formulations will be tested in vivo in mice to determine oral bioavailability of curcumin as a future diagnostic tool for AD.

EMPOWERING NURSES: USING KNOWLEDGE TO SCREEN & IDENTIFY VICTIMS OF HUMAN TRAFFICKING

P Khanna*, F Dong, R Trudgeon, IC Tuason. Western University of Health Sciences, Pomona, CA

Purpose of study Nursing professionals are indispensable to the care team when serving victims of human trafficking. Knowledge & trauma informed training are pivotal for developing & implementing screening tools for potential victims. Purpose is to assess the effectiveness of an educational intervention to increase the knowledge of current graduate nursing students about resources and federal laws for victims of human trafficking.

Methods used Pre-licensure students & licensed nursing students at Western University of Health Sciences were invited to participate in an educational activity about human trafficking. Participants were asked to fill out a pre seminar questionnaire. A 30-minute presentation was delivered. After the presentation, participants completed a post seminar questionnaire that consisted of the same ten questions from pre seminar questionnaire.

Summary of results Final analysis included 65 students. There was significant increase in awareness about human trafficking (84.9% vs 97.8%, p=0.0082). 7% corrected rank order of at risk victims before lecture and 15.2% post lecture (p=0.1025). Statistically significant change to correct response to questions was noted; which standard did US meet to eliminate human trafficking (64.8% vs 93.5%, p=0.0002), number of different types of human trafficking (16.3% vs 76.1%, p<0.0001), & what are the hotspot of human trafficking (20.4% vs 76.1%, p<0.0001).
Abstract 190 Table 1  Adjusted reimbursement trends, all values adjusted for inflation

<table>
<thead>
<tr>
<th></th>
<th></th>
<th></th>
</tr>
</thead>
<tbody>
<tr>
<td>66984</td>
<td>Removal of cataract with insertion of lens</td>
<td>-38.79%</td>
</tr>
<tr>
<td>66770</td>
<td>Injection of drug into eye</td>
<td>-59.14%</td>
</tr>
<tr>
<td>66821</td>
<td>Removal of recurrent external in lens capsule using laser</td>
<td>-22.82%</td>
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<tr>
<td>68761</td>
<td>Closure of tear duct opening using plug</td>
<td>-10.00%</td>
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<tr>
<td>68385</td>
<td>Laser repair to improve eye fluid</td>
<td>-47.43%</td>
</tr>
<tr>
<td>67820</td>
<td>Removal of cataract in lens using forceps</td>
<td>-44.25%</td>
</tr>
<tr>
<td>67190</td>
<td>Laser destruction of retinal growth</td>
<td>-32.65%</td>
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<tr>
<td>67904</td>
<td>Repair of tendon of upper eyelid</td>
<td>-16.45%</td>
</tr>
<tr>
<td>67228</td>
<td>Laser destruction of leaking retinal blood vessels</td>
<td>-69.30%</td>
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<tr>
<td>66761</td>
<td>Creation of eye fluid drainage tract</td>
<td>-35.55%</td>
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<tr>
<td>67840</td>
<td>Removal of cyclitic growth</td>
<td>3.84%</td>
</tr>
<tr>
<td>60800</td>
<td>Pulsing of nasal turbinates</td>
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<tr>
<td>67917</td>
<td>Repair of detached retina and drainage of eye fluid between the lens and retina</td>
<td>-31.42%</td>
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<tr>
<td>67145</td>
<td>Extensive repair of tearing-outward cyclitic defect</td>
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<td>66810</td>
<td>Dilatation of tear-drainage opening</td>
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<td>65800</td>
<td>Aspiration of eye fluid</td>
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<td>67515</td>
<td>Preventive retinal detachment treatment by heat or laser</td>
<td>-49.95%</td>
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<tr>
<td>67006</td>
<td>Removal of eye fluid (extreme) between the lens and retina</td>
<td>-26.75%</td>
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<td>67000</td>
<td>Repair of brow paralysis</td>
<td>-13.54%</td>
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<td>67810</td>
<td>Shunt of eyelid</td>
<td>-34.82%</td>
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<tr>
<td>Average</td>
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<td>-20.82%</td>
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</tbody>
</table>

Impact of Waiting Room Art Intervention on Pediatric Patients’ Anxiety and Cooperation with Physician

P Matus*, A Belikova, F Dong, S Fuchs. Western University of Health Sciences, Pomona, CA
10.1136/jim-2019-WMRC.191

Purpose of study  To determine whether the implementation of an art activity decreases pediatric patients’ anxiety in the waiting room and increases cooperation with their healthcare provider. This is a longitudinal pilot study that will expand on existing pediatric anxiety studies. The relationship between art therapy and pediatric stress levels has not been evaluated in detail in this specific circumstance.

Methods used  Research is being conducted in the Patient Care Center at Western University of Health Sciences, Pomona, CA, and is approved by the Institutional Review Board. A randomized controlled study design is done with pre-and-post intervention evaluations of both the intervention and control groups. The subjects in the experimental group perform a creative art activity while control subjects choose their own activity. Anxiety is evaluated when the subjects enter the waiting room and before the physician encounter using a modified SCARED questionnaire. The physician assesses the quality of the encounter by completing a questionnaire. Data will be analyzed using the SAS software for Windows version 9.3. Descriptive statistics will be presented as means and standard deviations for continuous variables, along with frequencies and proportions for categorical variables. The change in the anxiety score will be calculated as the summation of the five key items. The change will be assessed by the general linear model with the group (control vs intervention) as the categorical variable. All statistical analyses will be two-sided. P-value<0.05 will be considered to be statistically significant.

Summary of results  We are currently completing data collection. We hypothesize that an art intervention will reduce pediatric patients’ anxiety. This will result in better cooperation with the physician during the medical encounter. Data analysis will focus on quantifying subjects’ daily anxiety and situational anxiety in the PCC waiting room.

Conclusions  Conclusions are pending until an appropriate amount of data is collected. If our hypothesis is supported by this pilot study, we will expand this research to a larger number of subjects and facilities to obtain confirmation. This can yield recommendations for pediatric waiting room settings. *AB and PM have contributed equally to this work.
Conclusions According to our results, Medicare reimbursements have increased for longer patient visits and decreased for shorter patient visits. Additionally, Medicare reimbursement has increased for new patient visits and decreased for existing patient visits. Healthcare quality and access could be impacted by these trends as physicians are incentivized to accept new Medicare patients rather than follow-up visits. Future studies should be done to further examine reimbursement trends.

Purpose of study As the elderly population of the United States grows, so does the Medicare population and the need for transplant surgery. This study examines the Medicare reimbursement trends for transplant surgeries from 2000 and 2019.

Methods used The Centers for Medicare and Medicaid Services website was utilized to find the most commonly utilized transplant procedures using the CPY 2017 PUF data file and searching for the term ‘transplant’ in the HCPCS description. Using The Physician Fee Schedule Look-Up Tool, national reimbursement averages were calculated from 2000–2019 and data was analyzed. Averages were adjusted for inflation using the consumer price index.

Summary of results Medicare reimbursement for included procedures decreased from 2000 to 2019 (-19.85% or 0.99%/year). The largest decreases were seen in ‘Harvest of donor bone marrow for transplantation’ (-45.75%), ‘Transplant of tendon and muscle rerouting at lower leg or ankle’ (-43.19%), and ‘Removal of bladder and lymph nodes on both sides of pelvis with transplantation of ureters to small or large bowel with creation of urinary opening’ (-34.46%). There were only 3 procedures with increased reimbursement.

Conclusions Medicare is continually decreasing physician reimbursement rates for most transplant surgeries. Unfortunately, this potentially creates barriers to access these life-saving procedures for Medicare patients. This trend could also present many difficulties for surgeons and hospitals.
(35%), slips or trips (19%), and assaults (7%). Resulting effects included sprains or strains (40%), low back pain (25%) and abrasions or contusions (19%).

**Conclusions**
Reported injuries were fewer than expected by USBLS criteria. So were RD days according to an ACOEM-affiliated database, unless PT or specialty referral were prescribed. Whether these differences in RD days reflect severity of the injuries or other factors will require additional analysis.

Care of hospital-based healthcare workers rendered in OSCs staffed by Advanced Practice Providers generally yielded very good results.

**Abbreviations**
OSCs, onsite clinics; PT, physical therapy; RDD, restricted duty days.

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**HEALTH COACHING: BIOMETRICS WERE BETTER AFTER A YEAR**

1L Raymond*, 1M Slomba, 1CF Ford. 1Atrium Health, Charlotte, NC; 2University of North Carolina, Chapel Hill, NC

10.1136/jim-2019-WMRC.195

**Purpose of study**
To compare glycemic control and other biometrics over 1 year in 2 groups of wellness program participants (WPP) at a rural North Carolina workplace. Group A WPP had high-risk biometrics and were provided health coaching. Group B WPP had lower risks and got no coaching.

**Methods used**
Glycated hemoglobin (HbA1c) was measured by high performance liquid chromatography. Fasting blood glucose (FBG), total cholesterol (TC), low- and high-density lipoproteins (LDL, HDL) and triglycerides (TG) were measured by autoanalyzer in 94 Group A WPP (age 48 ± 10 SD, 62% male) and 390 Group B WPP (age 45 ± 12 SD, 41% male).

A certified health and wellness coach provided 2 to 4 counseling sessions to Group A WPP depending on risk factors: pre-diabetic and diabetic HbA1c > 5.6 and > 6.4, respectively, and improved diet, exercise, and weight control. WPP in both groups got letters with their blood test results requesting they share them with their physicians.

**Summary of results**
By group definitions, initial values of FBG, lipoproteins and blood pressures (table 1) were all much less salubrious in Group A than B (p=0.02–0.0001). A year later, between-group differences remained except for TC and LDL. HbA1c improved in Group A but glycemic control by FBG waned in Group B. TC and LDL also worsened in Group B. HDL and TG improved in both groups, with small but significant falls in diastolic blood pressure (BP).

**Conclusions**
Health coaching was accompanied by improved glycemic control not found in uncoached WPP. HDL and TG improved in both coached and non-coached groups, as did diastolic BP. The durability of these improvements requires further investigation.

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**EVALUATION OF THREE CYTOMEGALOVIRUS IGG LATERAL FLOW ASSAYS FOR DETERMINATION OF CYTOMEGALOVIRUS SEROSTATUS**

1FA All*, 1L Joncas-Schronce, 1G Pepper, 2RD Stapleton, 3GD Rubenfeld, 1M Boeckh, 1AL Imayev. 1University of Washington School of Medicine, Seattle, WA; 2University of Vermont, Burlington, VT; 3Sunnybrook Health Sciences Center, Toronto, ON, Canada

10.1136/jim-2019-WMRC.196

**Purpose of study**
Cytomegalovirus (CMV) serostatus is a major determinant of CMV reactivation risk. Current clinical serology assays are limited by slow turnaround time and need for either trained personnel and/or specialized equipment.

**Methods used**
We assessed the performance of three rapid lateral flow assays (LFA) for detection of CMV IgG antibodies compared to a primary reference CMV enzyme-linked immunosorbent assay (Zeus ELISA™ CMV IgG), in serum from 200 consecutive adults, performed clinically. A subset (N=60) of samples for which there was discordance between the

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**Abstract 194 Table 1**

<table>
<thead>
<tr>
<th></th>
<th>OSCs only (N=47)</th>
<th>OSCs + PT (N=6)</th>
<th>OSCs + Referral (N=15)</th>
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<tr>
<td>Actual RDD</td>
<td>Expected RDD</td>
<td>Actual RDD</td>
<td>Expected RDD</td>
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<tr>
<td>6.8±7.2</td>
<td>12.3±8.6</td>
<td>21.0±0.0</td>
<td>15.5±10.7</td>
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<td>p=0.001</td>
<td>p=0.008</td>
<td>p=0.0375</td>
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**Abstract 195 Table 1**

<table>
<thead>
<tr>
<th></th>
<th>Group A</th>
<th>Group B</th>
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</thead>
<tbody>
<tr>
<td></td>
<td>Initial</td>
<td>1-year follow-up</td>
</tr>
<tr>
<td>HbA1c:</td>
<td>8.7±1.8</td>
<td>7.6±1.4</td>
</tr>
<tr>
<td>FBG:</td>
<td>123±51</td>
<td>117±40</td>
</tr>
<tr>
<td>TC:</td>
<td>171±38</td>
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<td>LDL:</td>
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<td>HDL:</td>
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<td>47±13</td>
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<tr>
<td>TG:</td>
<td>193±105</td>
<td>171±99</td>
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<tr>
<td>BP, systolic:</td>
<td>137±15</td>
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<tr>
<td>BP, diastolic:</td>
<td>86±7</td>
<td>84±8</td>
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<td>Weight, pounds:</td>
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<td>245±54</td>
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</table>

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**Infectious diseases**

6:00 PM

Thursday, January 23, 2020

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**Poster session**

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**Abstracts**
primary reference assay and one or more of the LFAs was also tested by a second reference method: LIAISON DiaSorin CMV IgG. LFA testing was done by a single blinded research technician and followed by inter-operator agreement assessment in three independent, blinded, non-laboratory-trained personnel, using the LFA with the best performance characteristics.

Summary of results By the primary reference ELISA, 93 patients (46.5%) were seropositive, and 107 (53.5%) were seronegative, and there was 100% agreement (60/60) with the subset tested by the second reference assay. The sensitivity and specificity of the three LFAs (Healgen, Qoolabs read with Automated Reader, and nanoComposix), compared to the ELISA reference assay, were: 86%/83%, 99%/93%, and 57%/97%, respectively. For the QNow assay, the sensitivity and specificity were similar when interpreted by visual inspection with a UV wavelength flashlight (97%/97%). There was inter-operator agreement for 29/30 (97%) observations (10 samples; 4 positive, 6 negative).

Conclusions All three LFAs were rapid, easy to perform and interpret, but showed variability in performance. The ease of performance and rapid turnaround time provide the rationale to further evaluate these assays in clinical settings where these attributes would be advantageous.

Abstract 196 Table 1 Summary of diagnostic accuracy studies of the three lateral flow assays only 100 samples were used for the QNow UV Flashlight reader

<table>
<thead>
<tr>
<th>Lateral Flow Assays</th>
<th>Sensitivity (95% CI)</th>
<th>Specificity (95% CI)</th>
<th>PPV (95% CI)</th>
<th>NPV (95% CI)</th>
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</thead>
<tbody>
<tr>
<td>Healgen</td>
<td>86 (79 to 93)</td>
<td>83 (76 to 90)</td>
<td>82 (74 to 89)</td>
<td>87 (81 to 94)</td>
</tr>
<tr>
<td>QNow Automated Reader</td>
<td>99 (97 to 100)</td>
<td>93 (87 to 98)</td>
<td>93 (88 to 98)</td>
<td>99 (97 to 100)</td>
</tr>
<tr>
<td>nanoComposix</td>
<td>57 (47 to 67)</td>
<td>97 (94 to 100)</td>
<td>95 (89 to 100)</td>
<td>72 (65 to 80)</td>
</tr>
<tr>
<td>QNow UV Flashlight</td>
<td>97 (92 to 100)</td>
<td>97 (92 to 100)</td>
<td>95 (88 to 100)</td>
<td>98 (95 to 100)</td>
</tr>
</tbody>
</table>

197 COCCIDIOIDAL MENINGITIS AND CIRRHOSIS: IS THE GAME OVER?

1C D’Assumpcao*, 2FC Venter, 2K Sabatian, 2AA Heidari. 1Kern Medical – UCD, Bakersfield, CA; 2Valley Fever Institute, Bakersfield, CA

10.1136/jim-2019-WMRC.197

Introduction Coccidiodial meningitis requires lifelong, potentially hepatotoxic, triazole therapy. Complicated treatment cases might need neurosurgical shunt or reservoir for intrathecal agreement treatment. Cirrhosis complicates treatment options due to impaired hepatic function, coagulopathy and thrombocytopenia. We share a challenging case of coccidiodial meningitis in a cirrhotic patient with thrombocytopenia.

Case report 62 year old Hispanic male with alcoholic cirrhosis, portal hypertension requiring transjugular intrahepatic portosystemic shunt (TIPS), thrombocytopenia with splenic embolization, protein S deficiency with thrombosis of portal vein and left cephalic vein now off warfarin for two years, prediabetes and pulmonary coccidioidomycosis for two years off fluconazole for two months due to insurance presented with six weeks of progressive persistent occipital headaches, photophobia, nausea, vertigo, tinnitus, decrease hearing, blurry vision and short term memory lapses. Initial CT neuroimaging found 8 mm hyperdense focus in the posterior left cerebellum. Platelets were 51,000/uL. Presenting cirrhosis prognostic scores were Child Pugh A and MELD 10. TIPS was non-functional by ultrasound. MRI brain found mild hydrocephalus and abnormal periventricular signal in midbrain structures with severe leptomeningeal enhancements. Lumbar puncture had normal opening and closing pressures, lymphocytic pleocytosis, hypoglycorrachia, and elevated protein. Serum and CSF coccidioidal antibody immunodiffusion were reactive with severely elevated complement immunofixation titer. Due to thrombocytopenia intrathecal amphotericin was deferred. Because of fluconazole failure and voriconazole hepatotoxicity, isavuconazole was started. Dexamethasone taper was added due to hearing and short term memory loss. Outpatient audiogram and liver function monitoring was arranged with plan for lumbar puncture in four weeks. The pharmacokinetic and pharmacodynamic impact on his lifelong azole therapy from TIPS recanalization and impaired cytochrome P450 3A4 and 3A5 is unknown.

Conclusions Coccidiodial meningitis is fatal if not treated. Hepatic impairment complicates management due to therapeutic limitations. Prognostic impact needs to be studied further but, in our experience, it is diminished.

198 LYME DISEASE: DIVERSITY OF BORRELIA SPECIES IN CALIFORNIA AND MEXICO BASED ON A NOVEL IMMUNOBLOT ASSAY

1MC Fesler*, 2JS Shah, 3JJ Burrascano, 1RB Stricker. 1Union Square Medical Associates, San Francisco, CA; 2GeneX Laboratories, Milpitas, CA; 3Atkins Veterinary Services, Calgary, AB, Canada

10.1136/jim-2019-WMRC.198

Purpose of study Lyme disease is a growing tickborne epidemic with more than 400,000 new cases reported each year in the USA. If left undiagnosed and untreated, many of these cases progress to chronic Lyme disease, as recently defined (Stricker & Fesler, Am J Infect Dis 2018;14:1–44). Although Borrelia burgdorferi sensu stricto is considered the primary agent of Lyme disease in North America, recognition of previously undetected species of Borrelia burgdorferi sensu lato (Bbsl) and Relapsing Fever Borrelia (RFB) has complicated the diagnosis and treatment of Lyme disease. We report preliminary results of a serological survey of Bbsl and RFB in California and Mexico using a novel immunoblot technique.

Methods used The immunoblot method was designed to detect seroreactivity with specific species of Bbsl and RFB based on recombinant Borrelia membrane proteins, as previously described (Liu et al, Healthcare 2018;6:99; Shah et al, Healthcare 2019, in press). Patients were considered seropositive if they reacted with at least two proteins from a specific Borrelia species.
Summary of results Sixty-seven patients who resided in California or Mexico met the clinical definition of chronic Lyme disease. Immunoblot testing revealed that 30 patients were seropositive for Bbsl (Group 1), while 44 patients were seropositive for RFB (Group 2). Group 1 included patients who were seropositive for B. californiensis (6), B. spielmani (8), B. afzelii/B. garinii (8), and mixed infections that included B. mayonii (3). Group 2 included patients who were seropositive for B. hermsii (5), B. miyamotoi (7), B. turicatae (7) and B. turicatae (2). Seven patients were seropositive for both Bbsl and RFB species. In addition, four patients in Group 1 were seropositive for two species of Bbsl, while two patients in Group 2 were seropositive for two species of RFB. In the remaining Group 1 and Group 2 patients, the exact Borrelia species could not be identified using the immunoblot technique.

Conclusions Lyme disease is associated with a diversity of Borrelia species in California and Mexico. The clinical significance of this diversity in terms of missed diagnosis and response to antibiotic therapy requires further study.

Purpose of study Group C streptococci (GCS), specifically Streptococcus dysgalactiae subspecies equisimilis, are gram-positive, ß-hemolytic bacteria frequently isolated in the gastrointestinal tract, urogenital tract and occasionally as a part of the normal skin flora. We are presenting a complicated case of hip arthroplasty infection with Streptococcus dysgalactiae subspecies equisimilis.

Methods used Retrospective chart review.

Summary of results An 80-year-old female with a right total hip arthroplasty over 10 years ago was transferred to our hospital for fever, right hip swelling and pain. She stated that her hip dislocates every 6 to 8 months. Imagine showed underlying greater trochanter fracture and absent of hip abductors resulting in trochanteric escape and instability with hip dislocations. Imaging also revealed thick walled collection lateral to the hip extending to left iliacus muscle. She underwent incision and drainage and culture grew Streptococcus dysgalactiae subspecies equisimilis (SDSE). Sensitivity showed benzylpenicillin 0.32 mcg/mL and ceftriaxone 0.064 mcg. After 2 incision and drainage, She underwent 2 stage total hip revision. She continued on ceftriaxone 2 gram for 6 weeks.

Conclusion Streptococcus dysgalactiae subspecies equisimilis is an emerging bacterial infection to humans. Infection of Total Hip Arthroplasty with this pathogen is rare and has not been published. Streptococcus dysgalactiae, virulence, skin and soft tissue, infection, osteomyelitis, management.

REFERENCES
Acyclovir 200 mg, Ceftriaxone 2g and Vancomycin 320 mg. A lumbar puncture showed WBCs of 720, glucose of 27 and proteins of 128. CSF gram stain noted no organisms but with many PNMs. Patient continued to have nausea, emesis and fevers for 48 hours with temperatures up to 103. On day 3 of admission, serum coccidioidomycosis results indicated no IgM but a positive IgG with titers of 1:4 whereas the CSF results were negative. The patient was started on oral Fluconazole 200 mg and then transferred to a pediatric higher level of care. During transfer the patient suffered a grand-mal seizure for 1.5 minutes. He was then treated at the facility with Fluconazole 12 mg/kg/day and Keppra 30 mg/kg/day. Patient was discharged one week later with oral Fluconazole 100 mg twice a day and told to follow up for repeat LP and serum titers. An EEG conducted during his stay indicated mild encephalopathy. Patient would require Fluconazole as a lifetime treatment.  

**Conclusion** In the course of a coccidioidomycosis infection less than 50% have IgM present a week after symptom onset indicating a high false negative during the early stage of infection. The stated patient had no antibodies in the CSF. As a result, following up with a repeat LP is crucial to confirm the etiology of his meningitis. Very few cases of pediatric cocci meningitis have been reported. As a result, establishing a standard of care when encountering such cases is imperative.

**203 BACTERIAL TWO-COMPONENT SYSTEMS AS ANTIMICROBIAL TARGETS**


**Purpose of study** Two-component systems (TCS) comprising sensor histidine kinase (HK) and response regulator proteins are ubiquitous bacterial signal transduction mechanisms with regulatory functions in virulence, antibiotic resistance, toxin expression, cell differentiation and others. Some TCS are essential for bacterial viability yet no similar mechanisms are found in the animal kingdom. TCS are thus considered attractive antimicrobial targets. Previous efforts have identified lead compounds that inhibit multiple HK proteins and feature promising broad spectrum antimicrobial activities. Medicinal chemistry efforts resulting in a series of 50 novel compounds were probed for improved antimicrobial activities of the initial lead compounds.

**Methods used** Antimicrobial activities of 50 novel compounds against *Staphylococcus aureus* were assessed by Kirby-Bauer disk diffusion assays following Clinical & Laboratory Standards Institute guidelines. Minimal inhibitory concentrations (MICs) for the most promising compounds were assessed against a variety of Gram-negative and Gram-positive bacterial pathogens by serial dilution. All compounds were assayed for in vitro HK inhibitory activities and off-target inhibition of structurally related DNA-gyrase and structurally unrelated serine/threonine kinases were determined.

**Summary of results** Antimicrobial assays of the compound series singled out three with MICs in the µg/mL range.
against all tested bacterial pathogens. The compound series revealed inhibitors with varying specificities for a set of three HK in vitro, as assayed by autoradiography. None of the compounds showed off-target inhibition of DNA-gyrase, which shares a structurally related ATP-binding domain with HK proteins. Similarly, structurally unrelated serine/threonine kinases were not inhibited by any of the compounds.

**Conclusions** Novel compounds with promising antimicrobial activities were identified. TCS continue to be attractive targets and our efforts contribute towards a body of research with the ultimate goal of identifying effective inhibitors that can serve as new antibiotic agents.

**Poster session**

**Morphogenesis and malformations**

6:00 PM  
Thursday, January 23, 2020

**204 BILATERAL AGENSIS OF THE INTERNAL CAROTID ARTERIES**  
TS Matern, A Shinagawa*, R Rangasevamy, University of Nevada, Reno, Reno, NV  
10.1136/jim-2019-WMRC.204

**Introduction** Bilateral internal carotid artery agenesis (BLICAA) is a rare anomaly with minimal reporting in the literature. This case of BLICAA, consistent with previous case reports, presented with neurological symptoms.

**Case description** This case outlines an instance of bilateral ICA agenesis in a 59 year old male. The patient was hospitalized for acute onset left arm numbness. On admission, imaging revealed bilateral agenesis of the ICA. The intracranial vertebral arteries demonstrated atherosclerotic plaque with ~50% occlusion. MRI C-Spine also revealed severe left neuroforaminal stenosis at C6-7. The patient continued to experience left arm weakness and numbness, in a non-dermatomal distribution, with decreased bicep deep tendon reflexes. He denied associated radicular pain. The patient was discharged and instructed to follow-up with neurosurgery as an outpatient. Follow-up MRI revealed multiple lacunar infarcts. These imaging findings, in conjunction with an absence of radicular pain, were determined to be the cause of the patient’s left arm weakness and numbness.

**Discussion** This case is classified as an instance of BLICAA (delineated from aplasia by the absence of carotid canal on CT), of which only 32 cases were found to be previously reported in the literature. Furthermore, this patient presented with two additional vascular variations; first, a common origin of the right vertebral artery and common carotid artery, and second, an aberrant right subclavian artery. It is unclear exactly how this patient’s vascular anomalies contributed to his presentation, but in the context of previous cases, the absence of both ICAs was presumably a significant factor in the development of his cerebrovascular symptoms.

**Poster session**

**Neonatal pulmonary**

6:00 PM  
Thursday, January 23, 2020

**205 DOES CESAREAN SECTION INCREASE THE RISK OF LOWER RESPIRATORY TRACT INFECTIONS IN INFANTS?**  
1,2HV Dayag*, 2B Myint, 2KO k, 2V Sampath, 2J H ockman, 2A Yan, 2P Luu, 2,3B Afghani.  
1University of California, Irvine, Irvine, CA; 2UC Irvine School of Medicine, Irvine, CA; 3CHOC Hospital of Orange County, Orange, CA  
10.1136/jim-2019-WMRC.205

**Purpose of study** There has been some suggestion that infants born by C-section may be at higher risk of lower respiratory tract infections (LRTI) but the evidence is not clear. The objective of this study is to determine if c-section is associated with increased risk of LRTI in infants.

**Methods used** A literature review was conducted using Google Scholar and PubMed using key terms such as: lower respiratory tract infection (LRTI), c-section bronchiolitis, and pneumonia. Only studies that included infants born via c-section and a control group (vaginal delivery) with one or more LRTIs during the first 36 months of age were included in our analysis.
Abstract 205 Table 1  Association of c-section and lower respiratory tract infection in infants

<table>
<thead>
<tr>
<th>Author, Year; Location of Study</th>
<th>Type of Study</th>
<th>Average age to LRTI or Bronchiolitis</th>
<th>Outcome Details</th>
<th>N with LRTI/Total N C-section (C/S) Group</th>
<th>N with LRTI/Total N Vaginal Group</th>
<th>Statistical Significance</th>
<th>Confounding variables* accounted for (Y/N)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hendaus, 2014 - Qatar</td>
<td>Retrospective</td>
<td>10 days–34 months</td>
<td>Rate of RSV Hospitalization</td>
<td>89 (48%) of 185</td>
<td>215 (43%) of 493</td>
<td>p=0.965</td>
<td>N</td>
</tr>
<tr>
<td>Kristensen, 2015 - Denmark</td>
<td>Population-based cohort study</td>
<td>0–2 years</td>
<td>Rate of Hospitalization per1000 person-years for RSV</td>
<td>Acute C/S (N=31,715) 17.15 Elective C/S (N=30,965)=25.21</td>
<td>Vaginal (N=336,497) =15.56</td>
<td>Adjusted HR for RSV hospitalization = 1.09 (95% CI: 1.01–1.17, P=0.035) and 1.27 (95% CI: 1.19–1.36, P&lt;0.001), respectively.</td>
<td>Y</td>
</tr>
<tr>
<td>Magnus, 2011 - Norway</td>
<td>Prospective birth cohort study</td>
<td>0–36 months of age</td>
<td>Recurrent LRTI 3 or more episodes</td>
<td>Elective C/S 111 (6.6%) of 1,687, Acute C/S 143 (5.2%) of 2,736</td>
<td>1374 (4.4%) of 30829</td>
<td>Elective vs Vaginal Adjusted = 1.19, CI=(0.95–1.48) J Acute vs. Vaginal Adjusted RR=1.01, CI=(0.83–1.22)</td>
<td>Y</td>
</tr>
<tr>
<td>Moore, 2011 - Australia</td>
<td>Retrospective population-based cohort study</td>
<td>0–23 months</td>
<td>Hospitalization due to bronchiolitis</td>
<td>1051 (3.1%) of 33,421</td>
<td>3465 (2.7%) of 127,045</td>
<td>Adjustments for bronchiolitis at age = Y</td>
<td>Y</td>
</tr>
<tr>
<td>Shang, 2014 - China</td>
<td>Case-control study</td>
<td>less than 12 months</td>
<td>Respiratory Syncytial Virus, Bronchiolitis</td>
<td>104 (82%) of 126</td>
<td>69 (68%) of 101</td>
<td>Odds of RSV positive bronchiolitis was significantly higher in elective cesarean delivery (adjusted OR 2.32, 95% CI 1.19–4.52) after adjusting for confounding variables</td>
<td>Y</td>
</tr>
</tbody>
</table>

* The confounding variables included: maternal age, pregnancy complications, maternal smoking during pregnancy, maternal asthma. IHR: Hazard Ratio IRR: Incidence Rate Ratio

Summary of results We found 28 studies and of those, 5 met our inclusion criteria (see Table). Three of 4 studies that looked at the incidence of respiratory syncytial virus (RSV) showed a correlation between c-section and hospitalization due to RSV bronchiolitis. The correlation may be stronger in infants delivered by elective c-section vs emergent c-section (Shang et al). One study (Hendaus et al) did not find a correlation between c-section and recurrent LRTIs but this study did not take into account confounding variables, such as maternal age, pregnancy complications, maternal smoking and asthma.

Conclusions Our review suggests a possible link between c-section and increased risk of hospitalization due to RSV bronchiolitis. Large prospective studies that take into account various confounding variables are needed to confirm this association.

206 PRECISION CARE FOR ALVEOLAR CAPILLARY DYSPLASIA

CE Yost*, AR Putnam, L Jorgensen, RW Day. University of Utah, Salt Lake City, UT
10.1136/jim-2019-WMRC.206

Purpose of study The majority of patients with alveolar capillary dysplasia and misalignment of the pulmonary veins (ACD) die in early infancy from respiratory failure and pulmonary hypertension (PH). Here we report a patient with ACD who has survived for more than four years with precision care.

Methods used With IRB approval, we reviewed the medical records of a patient with ACD.

Summary of results Timeline: 0–3 months: The patient developed respiratory failure and pulmonary hypertension on the first day of life. He was refractory to treatment with high frequency ventilation, surfactant and inhaled nitric oxide (iNO); and was supported with ECMO from days of life 1–6. His inpatient therapy also included, epoprostenol, sildenafil and bosentan. Tests for genetic variants associated with surfactant deficiency were negative.

3–10 months: He was treated with sildenafil, bosentan, a diuretic and oxygen (O2). The severity of PH decreased and bosentan was stopped after 5 months.

11–24 months: He developed evidence of severe PH, right heart failure with an increase in B-type natriuretic peptide (BNP) and an increased O2 requirement with a respiratory infection. A CT angiogram raised concern for pulmonary veno-occlusive disease or pulmonary capillary hemangiomatosis. A lung biopsy showed evidence of a mild form of ACD. Genetic testing revealed a c.246C>G (p.F82L) heterozygous, novel and de novo variant in the FOXF1 gene. He was treated with bosentan, sildenafil, digoxin, aspirin and additional diuretics. His BNP normalized unless he experienced subsequent respiratory infections.

25–50 months: Heart catheterization and acute vasodilator testing (AVT) were performed to explore options for treatment with additional medications. His response to O2 with INO satisfied the Sitbon criteria for an acute responder. AVT with nicardipine confirmed that outpatient treatment with a calcium channel blocker would be appropriate. Amlodipine was added. Estimates of systolic pulmonary arterial pressure remain high; however, his BNP has been normal despite occasional respiratory infections and a decrease in diuretics.

Conclusions This case describes a patient with ACD who has survived for more than 4 years with medical therapy. His
acut response to nicardipine and his long-term response to amlopidine suggests that AVT can be used to guide precision care, even in a patient with right heart failure.

Poster session
Neonatology general
6:00 PM
Thursday, January 23, 2020

207 IMPROVING RESIDENT CONFIDENCE AT NEONATAL RESUSCITATION THROUGH INCREASED EXPOSURE

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10.1136/jim-2019-WMRC.207

Purpose of study Deterioration of resuscitation skills is seen with 39% of residents failing neonatal resuscitation program (NRP) skills shortly after completing training despite mandatory NRP training and adherence to ACGME requirements for newborn and NICU rotations. One study showed that pediatric residents were present at less than 50% of deliveries, declining to as low as 5%. The residents at our program attend very high risk resuscitations with a neonatologist, but are not required to attend normal term deliveries or lead resuscitations. This project hopes to advocate for increased exposure and a more deliberate curriculum toward resuscitative skills which has been shown to improve retention of NRP skills.

Methods used An initial anonymous survey was sent to all residents at our program (n=36) about their resuscitation experience. Normal newborn resuscitations (NNR) defined as a term newborn born via vaginal delivery with no pre-determined risk. A high risk delivery defined as any change from normal delivery, such as cesarean section but no neonatologist need be present. Responses (n=24) gave insight on (1) exposure to NNR (2) interest towards attending resuscitations (3) confidence in running resuscitations. Summary of results Data collection identified a total of 23 infants who met inclusion criteria, 13 infants were excluded, leaving 10 infants for analysis. Patient demographics included average BGA of 32w5d (23w4d–40w6d), 60% males, average BW of 2602 grams (565–4645 grams), and average CGA at time of initiation of ICM of 35w3d (25w1d–43w4d). Common primary diagnoses during ICM included pulmonary hypertension and hypoxic ischemic encephalopathy (HIE). Average exposure to ICM was 9.6 days (4–18.3 days). Average duration of the wean was 11.8 days (0–29 days). Infants weaned off directly from ICM compared to weaned off on a scheduled regimen had a shorter wean by 14.6 days (1.6 days versus 16.2 days).

Conclusions Our opioid weaning protocol has been approved by NICU physicians and pharmacists, and is now in the trial phase. The protocol will be implemented for infants on ICM, with the first cycle running from Nov-Dec 2019, and the second cycle Dec 2019-Jan 2020. After each cycle, we will assess not only duration of ICM weaning, but also adherence to the weaning protocol by practitioners and barriers to adherence.

208 EVALUATING AND IMPROVING CONTINUOUS MORPHINE WEANING PRACTICE IN THE NEONATAL INTENSIVE CARE UNIT

1A Iacob*, 1R. Meyer, 1A. Hayes. 1Miller Children’s and Women’s Hospital Long Beach, Long Beach, CA; 2Univ of California – Irvine, Orange, CA

10.1136/jim-2019-WMRC.208

Purpose of study In the Neonatal Intensive Care Unit (NICU), some critically ill infants require intravenous continuous morphine (ICM) for pain control, sedation, or comfort care. Exposure to ICM can lead to increased central line days, feeding intolerance, poor tone and motor function, and prolonged hospitalization. In our institution no protocol exists for weaning ICM which could contribute to longer morphine exposure than necessary. The goal of this Quality Improvement (QI) project is to shorten the ICM wean duration in our Level 3 NICU by 25% by Feb 2020 by creating a standardized protocol for weaning and discontinuation of ICM.

Methods used The first step was to analyze our current practice trends. All infants who received ICM for at least three consecutive days between Nov 2018 and Feb 2019 were evaluated for inclusion. Exclusion criteria were transfer to another facility or death while on ICM. The following data was collected for each patient: gestational age at birth (BGA), birth weight (BW), gender, gestational age (CGA) and primary diagnosis at initiation of ICM, duration of ICM, highest dose of ICM, duration of ICM wean, and duration and method of wean off ICM.

Summary of results The initial survey demonstrated that 92% of respondents requested more exposure to NNR in residency, which then led to a change in requirements. After implementing this change, there was an improvement in the number of residents who felt increased comfort at running resuscitations on their own from 21% to 58%. There was also improvement in knowledge regarding the type of equipment used at resuscitations.

Conclusions By increasing the number and variety of newborn resuscitations our residents are required to attend, our program has made their first steps in improving resident confidence and comfort in applying their NRP skills at neonatal resuscitations. This study implies that more exposure will continue to improve confidence in leading newborn resuscitations, and retention of NRP skills.
hypothermia (TH) and Veno-Arterial Extra Corporeal Membrane Oxygenation (VA-ECMO).

**Methods used** Single center retrospective evaluation of neonates with HIE on TH requiring VA-ECMO from 2011–2019.

**Summary of results** Among infants with HIE, meconium aspiration was the commonest cause of PPHN. All infants had echocardiographic evidence of PPHN and had received inhaled Nitric Oxide before ECMO. All except one infant tolerated complete course of TH for 72 h. Close monitoring of coagulation profile with prompt correction of abnormal values was performed during ECMO. All subjects were discharged home and 56% had normal MRI and only one patient had intracranial hemorrhage.

**Conclusions** Among patients with HIE, severe PPHN observed during TH can be effectively treated with VA-ECMO with overall good results. In patients with HIE and PPHN, TH through ECMO is an effective strategy.

### Abstract 209 Table 1 Neonatal characteristics and hospital outcomes - median (IQR) or n (%)

<table>
<thead>
<tr>
<th>Total patients</th>
<th>9</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age (weeks)</td>
<td>40 (39.3–40.3)</td>
</tr>
<tr>
<td>Male gender</td>
<td>3 (33.3%)</td>
</tr>
<tr>
<td>Birth weight (grams)</td>
<td>3480 (2820–3760)</td>
</tr>
<tr>
<td>Intubation in Delivery Room</td>
<td>7 (77.7%)</td>
</tr>
<tr>
<td>CPR in Delivery Room</td>
<td>6 (66.6%)</td>
</tr>
<tr>
<td>5 min Apgar score ≤ 5</td>
<td>7 (77.7%)</td>
</tr>
<tr>
<td>Blood gas for which cooling was initiated</td>
<td>- pH</td>
</tr>
<tr>
<td>- Base deficit - mmol/Ltr</td>
<td>17 (14–19)</td>
</tr>
<tr>
<td>Seizures</td>
<td>2 (22.2%)</td>
</tr>
<tr>
<td>Co-morbidities</td>
<td>- Meechanism</td>
</tr>
<tr>
<td>- Pulmonary hemorrhage</td>
<td>1 (11%)</td>
</tr>
<tr>
<td>- Trisomy 21</td>
<td>2 (22%)</td>
</tr>
<tr>
<td>- Sepsis</td>
<td>1 (11%)</td>
</tr>
<tr>
<td>Age at starting Nitric oxide (hrs)</td>
<td>4 (3–6)</td>
</tr>
<tr>
<td>Echo finding before ECMO</td>
<td>- Pulmonary hypertension</td>
</tr>
<tr>
<td>- Ventricular dysfunction</td>
<td>3 (33%)</td>
</tr>
<tr>
<td>Values just before ECMO</td>
<td>- Platelet count</td>
</tr>
<tr>
<td>- INR</td>
<td>1.47 (1.32–2)</td>
</tr>
<tr>
<td>- Fibrinogen</td>
<td>200 (176–225)</td>
</tr>
<tr>
<td>- Intracranial Hemorrhage</td>
<td>0</td>
</tr>
<tr>
<td>- Oxygenation Index</td>
<td>51 (43.5–60)</td>
</tr>
<tr>
<td>- Vasoactive Inotrope Score</td>
<td>22 (15–55)</td>
</tr>
</tbody>
</table>

**Characteristics during ECMO**

- Age at initiation (hrs of life) | 24 (22–36) |
- Cooling during ECMO* | 7 (77.7%) |
- Intracranial hemorrhage | 1 (11.1%) |
- Lowest PaCO2** | 27 (19–31) |
- Total ECMO duration (days) | 5 (4–7) |
- Duration of Mechanical Ventilation (days) | 15 (13–18) |
- Age at MRI (days) | 13 (10–15) |
- Normal MRI | 5 (55%) |
- Length of stay (days) | 36 (30–59) |

* One patient was rewarmed at referral hospital before transfer at 32 h of life for worsening PHTN. Second patient ECMO was started after 72 h of TH. ** Patient with INH had lowest PaCO2 of 17 mm Hg.

**Purpose of study** An operational definition neonatal sepsis is lacking. Technological tools and biomarkers that can accurately predict infection mortality outcomes, especially risks for multi-organ dysfunction syndrome, are constantly being sought. The neonatal sequential organ failure assessment (nSOFA) score, recently published by Dr. Wynn and Dr. Polin in Pediatric Research, provides an objective, electronic health record-automated score. Conversely, HeRO incorporates proprietary algorithms to calculate a sepsis score based upon variations in the patient’s heart rate. This study will independently validate the nSOFA score and directly compare the efficacy of nSOFA and HeRO scoring systems in predicting mortality related to neonatal sepsis.

**Methods used** A retrospective chart review of infants born <33 weeks’ gestational age between 2012 and 2015 at UCSD Medical Center with positive blood cultures and/or necrotizing enterocolitis (NEC) were included. The electronic health records of these infants were reviewed to ascertain and document components of the nSOFA score (range 0–15) including: (a) need for mechanical ventilation and oxygen requirement (score range 0–8); (b) administration of vasoactive drugs including corticosteroids (score range 0–4); and (c) presence and degree of thrombocytopenia (score range 0–3). Scores were calculated for each infant at nine time points, including −48, −24, −12, −6, time 0 (sepsis evaluation), +6, +12, +24, and +48 hours. nSOFA data was directly compared to HeRO scores at similar time point to determine efficacy in predicting neonatal mortality associated with sepsis and/or NEC.

**Summary of results** Seventy-five patients born <33 weeks’ gestational age were identified to have one or more positive blood cultures during the study period and were included for analysis. In addition, seventeen patients diagnosed with necrotizing enterocolitis during this time period have been identified. Chart review is nearly completed with final statistical analysis expected in November 2019.

**Conclusions** Complete results with conclusion will be available by the WSPR conference date.

**210 INDEPENDENT EVALUATION OF THE NEONATAL SEQUENTIAL ORGAN FAILURE ASSESSMENT SCORE WITH DIRECT COMPARISON TO HERO TECHNOLOGY**

1. B Barret, 2 JL Wynn, 3 S Lawrence*. 1UCSD, San Diego, CA; 2University of Florida, Gainesville, FL

**Purpose of study** Helping Babies Breathe (HBB) is an educational module created by the American Academy of Pediatrics that teaches the knowledge and skills required to care for babies in the first few minutes of life after birth. Since its creation in 2010, HBB has been implemented in 80 countries and has been taught to over 450,000 birth attendants resulting in a reduction of neonatal mortality by as high as 50%. The program relies on in-person trainings that cover both the knowledge and skills required to care for newborns. However,
recent studies have shown that low-dose high-frequency refresher trainings are an effective means of accomplishing this, but to date these trainings have not incorporated multimedia tools. While multimedia materials have already been created to aid in teaching HBB, their effectiveness has never been studied. This project aims to study the effectiveness of targeted low-dose high-frequency refresher videos of HBB material in one hospital in Tanzania.

Methods used Previously trained birth attendants and health care providers at Muhimbili National Hospital in Dar es Salaam, Tanzania will be recruited to participate in this study. Participants will be randomized to either a cohort that only views a refresher video, or a cohort that receives the standard in-person refresher teachings. Skills over time will be measured using the already established HBB bag and mask ventilation skills checklist. To establish a baseline, all participants will be scored by this checklist at the beginning of the study. Before the interventions are implemented, skills will be checked after 2 months to examine how skills change over time. Then, either a short refresher video or a hands-on person-to-person skills refresher will be presented to the respective cohorts. Skills stations will be set up in the labor and delivery area and all participants will self-practice bag and mask ventilation weekly for 5 minutes. The video cohort will be required to re-watch the skills video twice a month. The bag and mask ventilation skills checklist will be administered again after 2 months and the performance of the two cohorts will be compared.

Summary of results Pending

Conclusions It is hypothesized that the two cohorts will perform similarly which may suggest that self-study videos can replace in-person refresher trainings when trainers are not available.
Spontaneous bilateral subcapsular renal hematoma is a rare condition. Upon literature review, only two case reports have elucidated possible etiologies for such a presentation. Similarly, we present a 52-year-old female with PMH of DM type II, HTN, decreased cognitive function s/p TBI from MVA x2 last one in 2011, who developed spontaneous bilateral renal subcapsular hematoma on admission for management of DKA and pyelonephritis. Typically, subcapsular hematoma achieves spontaneous resolution with conservative management, however, our patient was having active bleed with hypotensive shock needing left renal artery embolization and eventually left nephrectomy. Her condition was further complicated with total left lung collapse secondary to mucous plug and septic shock.

**Conclusion**

The main interest of this case report lies in bilateral spontaneous subcapsular hematoma in a patient who has multiple factors that could have played a role in this condition. Our patient was treated with embolization, bilateral nephrostomy tubes, and nephrectomy of the left kidney. We present the first case of spontaneous bilateral subcapsular hematoma in the setting of no major risk factors that could possibly affect both kidneys simultaneously.

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**Purpose of study**

The operating room can be saturated with noxious smells. Anecdotally, medical staff apply products to lessen the impact of these smells. This study aimed to determine the odour-masking ability of four inexpensive and convenient products.

**Methods used**

This study was a randomized, single-blinded crossover study. Individuals between ages 19 and 30 years were included. Exclusion criteria included active allergies, upper respiratory tract infection, alteration to sense of smell, or failure of olfactory screen. Eighty-one individuals were recruited; one was excluded following a failed olfactory function test. Participants were exposed to an experimental odour in lieu of a noxious surgical odour. After smelling the experimental odour without barriers, participants were re-exposed to the odour using five surgical masks in randomized order. Each mask was lined with a test product (cherry lip balm, tincture of benzoin, Mastisol®, mint toothpaste, and control (plain mask)). Participants rated the effectiveness of products at masking the experimental odour from 0–100 (0 = completely ineffective; 100 = completely effective). Participants also rated the perceived effectiveness of each product overall. The main outcome measure was the odour-masking effectiveness of the four products.
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Summary of results Eighty participants were included in the study (33 male, 47 female), averaging 24.2 years of age. Mean odour-masking effectiveness for cherry lip balm was 66.5 (±24.6), tincture of benzoin: 62.6 (±25.0), Mastisol®: 61.3 (±23.9), mint toothpaste: 57.5 (±27.4), and control: 21.9 (±21.8). All products performed better than the control (p-value: <0.001), but there was no significant difference in performance between products. Cherry lip balm was the most preferred odour-masking product (29 participants), followed by mint toothpaste (22), Mastisol® (14), tincture of benzoin (10), and control (5).

Conclusions All tested products demonstrated odour-masking abilities. We recommend that healthcare professionals find the odour-masking product that works best for them, starting with cherry lip balm.

216 POINT OF CARE ULTRASOUND EVALUATION OF AIRSPACE DISEASE BEFORE AND AFTER PROLONGED SURGERY
K Cook*, M Buell, B Lew, P Vu, J Hinson, M Martinez, M Alschuler, M Holclaw, D Ramingih. Loma Linda University, Loma Linda, CA
10.1136/jim-2019-WMRC.216

Purpose of study Point-of-care ultrasound (POCUS) has rapidly emerged as a modality to improve bedside assessment of pulmonary air-space disease. Indeed, POCUS has demonstrated to be superior for the assessment of air-space disease in comparison to both chest x-ray and stethoscope auscultation. Given that patients undergoing a long surgical procedure requiring general anesthesia are at risk for pulmonary complications the ability to assess air-space disease perioperatively is of interest. The authors designed a feasibility study to evaluate the utility of point of care ultrasound to evaluate the development of air-space in this patient population.

Methods used Adult patients were screened for those undergoing a surgical procedure scheduled for more than 3 hours of surgical time. Patients were then scanned using a validated POCUS examination to detect both the presence and severity of air-space disease across 5 lung fields for each side (total of 10 views). POCUS exams were performed immediately before and after their surgical procedure. Airspace disease was defined by the presence of B-lines, as previously validated. Before-to-after comparisons in air-space disease was performed by Chi-Square Analysis in the frequency of B-Lines in each lung quadrant and in total. A p value of less than 0.05 was identified as significant.

Summary of results Preliminary data on 15 patients with 150 ultrasound images demonstrates significantly higher number of B lines post-surgery (17%) compared to pre-surgery (10%), p<0.05. The dependent lung zones were the most frequent areas to develop B lines both in the pre-op and post-op settings. All patients were able to have ultrasound images of the lung parenchyma across all lung fields.

Conclusions POCUS examination of the lungs fields can be successful performed in the perioperative setting. Our preliminary data supports this tool be useful in the assessment of the change in the aeration of lung tissue from pre to post surgery. Specifically, for patients undergoing long surgical procedures, this modality may be useful to detect the development of new areas of airspace disease.

218 ADULT MESENCHYMAL HAMARTOMA OF THE LIVER: CASE REPORT AND LITERATURE REVIEW
1M Gross*, 2N Wilkinson. 1University of Washington School of Medicine, Seattle, WA; 2Kalispell Regional Healthcare Surgical Specialists, Kalispell, MT
10.1136/jim-2019-WMRC.218

Case report A 39-year-old woman presented to the emergency department for acute onset epigastric pain. Imaging demonstrated a 5 x 6 cm lesion in segment 4 of the liver. A left hepatic resection was performed and the patient did well post operatively. Final microscopic diagnosis demonstrated mesenchymal hamartoma. Mesenchymal hamartoma of the liver (MHL) is a benign lesion, commonly seen in children younger than five years. MHL rarely occurs in adults, with only 31
Abstract 218 Figure 1 MRI of the abdomen with contrast demonstrating multiple masses in liver, with largest occurring within the anterior left hepatic lobe measuring 5.9 cm

cases reported worldwide as of 2010. The pathogenesis of MHL is not completely understood, though it is believed to relate to aberrant development of the bile ducts. Despite a typically benign course, malignant transformation may occur. Recent cytogenic analysis has identified chromosomal rearrangements nearly identical to those observed in undifferentiated embryonal carcinoma (UEC), an aggressive lesion with a median survival of less than 1.5 years. UEC has arisen from MHL in multiple instances, while hepatic angiosarcoma has also been reported. Additionally, MHL may grow to massive proportions. Due to the potential for such complications, aggressive surgical management with complete resection is mandated when feasible. Herein we discuss the aforementioned case as well as the epidemiology, clinical presentation, histopathology, cytogenetics, and treatment of this rare adult neoplasm.

Purpose of study Treatment resistant epilepsy is a major health burden. Recurrent seizures can result in negative sequela which include seizure related injury, cognitive and memory impairment, and increased healthcare utilization. Hemispheric surgery is an excellent and potentially curative option for recurrent seizures which include seizure related injury, cognitive and memory impairment, and increased healthcare utilization. Hemispheric surgery is an excellent and potentially curative option for recurrent seizures. To test this hypothesis, the medical records of all patients who had undergone hemispheric surgery at BC Children’s Hospital were evaluated. Data was gathered on clinical history, neuromaging, EEG reports, seizure types, seizure frequency pre- and post-surgery and AED use.

Summary of results We observed that over 90% of patients had very good outcomes in terms of seizure freedom with surgery but 43% percent of patients were still using AEDs post-operatively despite seizure freedom. Analysis of the data showed trends towards increased seizure recurrence and AED use with bilateral or generalized brain abnormalities on EEG, as hypothesized. However, these results were not significant, likely due to small sample size (n=40).

Conclusions Given the high success rate of hemispheric surgery, when seizure freedom is the metric, any modest prediction capability conferred by EEG analysis would be of limited clinical use. However, there may be a role for using pre-operative EEGs to inform medical care after surgery with regards to AED requirements and weaning.

Abstract 219 UTILITITY OF EEG IN PREDICTING HEMISPHERIC SURGERY OUTCOME

K Hanjani*, 1M Connolly. 1University of British Columbia, Vancouver, BC, Canada; 2BC Children’s Hospital, Vancouver, BC, Canada

Purpose of study Treatment resistant epilepsy is a major health burden. Recurrent seizures can result in negative sequela which include seizure related injury, cognitive and memory impairment, and increased healthcare utilization. Hemispheric surgery is an excellent and potentially curative option for carefully selected patients and is currently underutilized. One of the important modalities used to assess patient suitability for the surgery is the electroencephalogram (EEG), which can non-invasively study brain activity. The goal of this project was to evaluate the role of EEG in predicting surgery outcome.

Methods used We hypothesized that generalized or bilateral epileptic abnormalities captured on EEG would be associated with risk of ongoing seizures and ongoing need for antiepileptic drugs (AEDs) post-surgery. To test this hypothesis, the racial differences in the prostate cancer microenvironment.

Purpose of study African-American men (AAM) are more likely to be diagnosed with prostate cancer (PC), and have more aggressive PC at diagnosis, than any other ethnic group. Differences in inflammatory cell types among races may account for this imbalance. We previously found that AAM have more T-cells in their PC as compared to European American men (EAM). Herein, we developed a panel to quantify myeloid cells in the same tissue samples. We hypothesize there are racial differences in myeloid-derived suppressor cell (MDSC) counts in the prostate cancer microenvironment.

Methods used Using 5-plex immunofluorescence (IF) on prostate biopsy formalin-fixed paraffin-embedded (FFPE) slides of AAM and EAM matched controls and PC cases, we quantified granulocytic (G) and monocytic (M) MDSCs. The panel includes antibodies for CXCR2, CD14, CD11b, HLA-DR, and cytokeratin, enabling us to quantify the recruitment and density of M-MDSCs and G-MDSCs to the tumor microenvironment. M-MDSCs are defined as CD14+ and HLA-DR low/-, whereas G-MDSCs are defined as CD14+, HLA-DR low/-, and CD11b+. Each cell type was counted in PC regions as well as in biopsy negative controls. Data were then compared using unpaired t-tests with a=0.05/n, where n is the number of markers used, to accommodate multiple testing.

Summary of results Among all men with PC, we found higher M-MDSC density in cancers vs. controls, though these differences were not statistically significant (p=0.076). AAM PCs had significantly more M-MDSCs (p<0.017) and G-MDSCs (p<0.19) than AAM non-PCs, though the latter was not statistically significant. There was a trend for higher G-MDSCs in EAM non-PC tissue as compared to AAM non-PCs (p<0.055). AAM PCs had significantly more CXCR2+ cells than EAM PCs (p<0.008).

Conclusions We found racial differences in MDSC counts both in PC and non-PC prostate tissues. These preliminary results
Abstracts

221 COMPARATIVE ASSESSMENT OF PATIENT SURGICAL RISK BY SURGEONS VS. A UNIVERSAL, PARSIMONIOUS STATISTICAL RISK SYSTEM

N Mason*, P Razeboom, MR Bronsert, W Henderson, N Pradhan, A Lambert-Kiener, K Hammermeister, R Meguid. University of Colorado School of Medicine, Aurora, CO

10.1136/jim-2019-WMRC.221

Purpose of study The literature lacks evidence on accurate preoperative prediction of postoperative surgical outcomes by surgeons. The Surgical Risk Preoperative Assessment System (SURPAS) provides individual patients with accurate procedure-specific preoperative risk prediction of 30-day postoperative adverse outcomes including mortality, overall morbidity, & 9 other surgical complications. To predict these values, SURPAS uses 8 variables including procedural complexity & procedure-specific risk (both derived from the current procedural terminology code), functional health status, American Society of Anesthesiologists Physical Status Classification (ASA class), patient age, emergency status of the operation, in-/out-patient procedure, & surgeon specialty. These risk algorithms were developed from American College of Surgeons National Surgery Quality Improvement Program (NSQIP) data, which include the independent variables entered into SURPAS & the postoperative adverse outcomes.

Methods used We compared the accuracy of surgeons’ ability to predict overall morbidity & mortality for a variety of surgical procedures within their specialty to the outcomes predicted by SURPAS, & to the known postoperative outcomes. 30 patients’ NSQIP data was presented to surgeons in standardized vignette formats, including the procedure performed & field of expertise. We will measure surgeon accuracy.

Summary of results Preliminary results of demographic data showed no statistically significant difference between the two groups. (Table 1) Outcomes data showed a statistically significant increase in SICU and hospital LOS for PACU admitted patients. Mortality was also statistically significant but was higher for SICU admitted patients. (Table 2)

Conclusions We saw a median PACU LOS of 103 mins yet the PACU to SICU group’s SICU LOS was 844 mins longer than the PACU group. When it came to hospital LOS the PACU to SICU group had a 422 mins longer stay. Delay in ICU care leads to increased LOS in the SICU and the hospital. The mortality was significantly higher in the directly admitted SICU patients which from our preliminary analysis appears to be from patients who had high ASA and APACHE II scores and died soon after ICU admission.

222 EFFECTS OF DELAYED ICU ADMISSION FROM THE POST-ANESTHESIA CARE UNIT VS. DIRECT ADMISSION TO THE SURGICAL INTENSIVE CARE UNIT IN POSTOPERATIVE PATIENTS

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10.1136/jim-2019-WMRC.222

Purpose of study The post-anesthesia care unit (PACU) was established to minimize complications immediately after anesthesia and surgery. Often patients are held up in PACU due to bed availability or lack of appropriate staffing. This study was conducted with the hypothesis that PACU admission of a critically ill patient increases ICU length of stay (LOS). Secondarily we also looked at hospital LOS and mortality.

Methods used A retrospective chart review was conducted at a tertiary academic medical center between 2/1/13–6/1/19. We identified all surgical patients admitted to the SICU, a total of 5365 patients. We separated these patients into two groups: (1) direct SICU admission and (2) PACU then SICU admission. We collected the following data: (1) demographic characteristics; (2) the American Society of Anesthesiologists (ASA) score; (3) PACU LOS; (6) SICU LOS; (7) Hospital LOS and (8) mortality.

Summary of results Preliminary results of demographic data showed no statistically significant difference between the two groups. (Table 1) Outcomes data showed a statistically significant increase in SICU and hospital LOS for PACU admitted patients. Mortality was also statistically significant but was higher for SICU admitted patients. (Table 2)

Conclusions We saw a median PACU LOS of 103 mins yet the PACU to SICU group’s SICU LOS was 844 mins longer than the PACU group. When it came to hospital LOS the PACU to SICU group had a 422 mins longer stay. Delay in ICU care leads to increased LOS in the SICU and the hospital. The mortality was significantly higher in the directly admitted SICU patients which from our preliminary analysis appears to be from patients who had high ASA and APACHE II scores and died soon after ICU admission.
SURGICAL OUTCOMES OF SUPRAVALVULAR AORTIC STENOSIS REPAIR TECHNIQUES IN PEDIATRIC PATIENTS: A SYSTEMATIC REVIEW

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10.1136/jim-2019-WMRC.223

Purpose of study Supravalvular aortic stenosis (SVAS) is a congenital anomaly arising from familial or sporadic mutations in the elastin gene. This mutation results in severe constriction of the sinotubular junction. SVAS is most commonly seen as a feature of Williams Syndrome. One, two, and three patch aortoplasty techniques are the most frequently performed corrective procedures for SVAS worldwide and have shown efficacious outcomes in the repair of SVAS. However, data comparing the outcomes and survival rates between these techniques are limited.

Methods used A systematic literature review using PubMed and EMBASE identified 15 papers for data extraction related to pediatric SVAS surgical outcomes and survival. Papers were selected through systematic use of search terms and were manually assessed for relevance and inclusion of pertinent data. These data included re-operation rates, pre/post-operative Doppler gradients, mortality, and significant comorbidities. Review and meta-analysis were conducted according to PRISMA guidelines.

Summary of results The average age of pediatric patients assessed was 7.8 years ± 3.2 with 44% of patients presenting with Williams syndrome. The pre/post-operative Doppler gradients showed no significant difference between the one, two, and three patch aortoplasty techniques. Similarly, early and late mortality was not significantly different between techniques. However, reoperation rates due to restenosis varied with an incidence of reoperation of 13.8%, 4.8%, and 0% in patients with one, two, and three patch aortoplasties, respectively.

Conclusions In pediatric SVAS patients, one, two, and three patch aortoplasty techniques show no significant differences in pre/post-operative gradients and survival rates, while three patch aortoplasty reveals decreased incidence of reoperation. Further studies are needed to better delineate the best surgical approach.

OBSTRUCTIVE SLEEP APNEA AND EARLY WEIGHT LOSS AMONG ADOLESCENTS THAT UNDERGO BARIATRIC SURGERY AT CHILDREN’S HOSPITAL COLORADO

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10.1136/jim-2019-WMRC.224

Purpose of study Adult literature demonstrates impressive results regarding the effect of bariatric surgery in ameliorating obstructive sleep apnea (OSA) however data are scarce in pediatric literature. As morbid obesity and OSA become increasingly common in adolescents, it is important to quantify the prevalence of OSA and the effects of surgical intervention in this unique, growing population.

Methods used Retrospective chart review of adolescents enrolled in the Bariatric Surgery Center at Children’s Hospital Colorado (CHCO) with pre- and post-surgical polysomnography (PSG) between 06/17–08/19 (70/83 total patients). Inclusion criteria was based on availability of presurgical PSG results. Variables including age, gender, race, comorbidities, PSG results, and weight were collected. Pediatric OSA criteria were applied to PSG results to determine presence of OSA. Tests for nonparametric data were used to compare baseline characteristics and surgical outcomes between patients with and without preoperative OSA.

Summary of results The prevalence of OSA among those with preoperative PSG, defined as apnea hypopnea index ≥1, was 77% pre-surgery, with 44% demonstrating severe OSA by OAH ≥10. There were no significant differences in race, gender, BMI, or comorbidities according to diagnosis of OSA. Of 12 patients with preoperative OSA and a postoperative sleep study, 38% resolved their OSA an average of 4.92 months post-surgery. Average preoperative BMI for resolving patients was 45.46 versus 56.76 in those who did not have resolution of OSA (p=0.03). Average change in BMI from pre- to post-operative sleep study was -21.9% in the resolved group and -13.5% in the unresolved group (p=0.12).

Conclusions The prevalence of OSA in the population of adolescents seen in the Bariatric Surgery Center at CHCO was more similar to the estimated prevalence in adults seeking bariatric surgery (74%) than the general population of adolescent with obesity (43%). Resolution of OSA after surgery correlated with lower preoperative BMI, but sample size limited further analysis.

CLINICAL OUTCOME DIFFERENCES BETWEEN SINGLE AND MULTI-STAGED TRANSTIBIAL AMPUTATIONS

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Purpose of study Transtibial amputations are currently performed as single guillotine amputations or multi-staged amputations with primary and formalization amputations. This study assessed if multi-staged amputations provide improved clinical outcomes over single stage transtibial amputations and what patient groups it may benefit.

Methods used We queried 207 records of patients receiving transtibial amputations from January 2015 through December 2018. Patient demographics, comorbidities, preoperative factors, surgical factors, and complications were recorded. Regression modeling was utilized to compare functional outcomes and influential perioperative variables between single and multi-staged amputation groups. After accounting for exclusion criteria, 148 patient records were analyzed.

Summary of results From the records analyzed, 49 patients (37%) underwent multi-stage amputation. Patients with multi-stage amputations had a hazard ratio of 0.60 for complication risk (P=0.04). The most common complications, often requiring readmission, were infection (n=30) and wound dehiscence (n=21). This was the case for both single and multi-staged amputations. For patients who were readmitted, 72% required amputation revision. Of the patients who experienced complications, 56% of patients were independently ambulatory with a prosthesis and 31% were ambulatory with additional assistive devices (e.g., cane, front
SLEEP BEHAVIORS IN ADOLESCENTS UNDERGOING BARIATRIC SURGERY EVALUATION

Purpose of study Poor sleep behaviors has been associated with increased risk of obesity and insulin resistance. Sleep health has not been examined in adolescents with obesity undergoing bariatric surgery. Methods used A chart review was performed and all patients receiving care at the Children’s Hospital Colorado (CHCO) Bariatric Surgery Center between 06/17–08/19 were included. Demographic, medical history, sleep behaviors, and laboratory measures were abstracted. T-tests were used to examine the differences between sleep variables and baseline weight and comorbidity status. Summary of results Data from 85 patients, aged 16.8 years, were identified. The majority of patients were female (71%) and Hispanic (52%). The majority of patients reported a family history of obesity (69%) and T2D (80%). Over half (52%) reported receiving mental health counseling previously. Pre-surgery, 24% of the patients were diagnosed with hypertension and 20% with Type 2 Diabetes. A total of 29% screened positively for anxiety and 41% were diagnosed with depression. Over half of the patients (60%) stated they did not have a regular sleep schedule. A typical total time in bed of 9 hours was reported with as much as 86 minutes of variability. A total of 36% of patients reported a bedtime of midnight or later and 78% were documented to have a mobile device with them in their bed. Patients who reported variability in their sleep schedule of at least 60 minutes were found to have significantly higher weights (139.1kg vs 131.1kg; p=0.02) and BMI (48.8 vs 44.9; p=0.04) at baseline. Similarly, patients who reported less than 8.5 hours of time in bed each night had significantly higher weights (144.7kg vs 131.7kg; p=0.002), BMI (50.3 vs 45.9; p=0.02), and HbA1c levels (6.18 vs 5.85; p<0.001), compared to patients that had greater than 8.5 hours of time in bed each night. Conclusions In a population of adolescents seen in the bariatric surgery center at CHCO, sleep behaviors preoperatively were related to baseline weight, BMI, and HbA1c levels. Insufficient sleep and unhealthy sleep behaviors may adversely impact adolescents’ overall health. Interventions to improve sleep health should be evaluated as part of efforts to improve health outcomes in patients undergoing bariatric surgery.

OPTIMAL NAIL DIAMETER TO MEDULLARY CANAL RATIO IN DIAPHYSIAL TIBIA FRACTURES TREATED WITH INTRAMEDULLARY NAILING

Purpose of study Up to 17% of diaphyseal tibia fractures result in delayed union. Most patients achieve excellent outcomes with intramedullary nailing and it is unknown why some experience delayed healing. The goal of our study was to assess potential risk factors that may influence fracture healing in these patients with an emphasis on the nail diameter to medullary canal (ND/MCD) ratio. Methods used Adult patients that underwent intramedullary nailing of tibia fractures over a 10-year period were retrospectively reviewed. Exclusion criteria were inadequate follow-up (<12 months), additional lower extremity fractures, additional hardware, non-diaphyseal and pathologic fractures. Post-operative anteroposterior (AP) and lateral radiographs were used to calculate the ND/MCD ratio. Multi-variable logistic regression analyses were used to identify demographic and clinical variables associated with complications. A receiver operating curve analysis was used to identify the ND/MCD ratio that best differentiated between subjects who developed a nonunion and those who did not. Summary of results The average age among the 95 individuals included in the study was 44.2 years (±16.1). The cumulative incidence of complications was 29% [95% CI: 20.3 to 39.7%]. The presence of an open fracture was the only variable significantly related to the risk of a complication. The odds of a complication among open fractures were 10.1 times [95% CI: 3.2 to 32.1, p<0.0001] times the odds of a complication among closed fractures. Nonunion was noted in 18 patients [19.0%, 95% CI: 11.1 to 26.8%]. Age, sex, ND/MCD ratio, and presence of an open fracture were used to build a logistic model to predict nonunion (AUC=0.83; 95% CI: 0.71 to 0.96). ND/MCD ratio cutoff of 85% was associated with the highest AUC value (sensitivity=44%, specificity=79%) in an exploratory analysis differentiating between the subjects that developed nonunion and the ones that did not. Conclusions Presence of an open fracture is strongly associated with increased complication rates among patients undergoing intramedullary nailing of tibia fractures. Canal fill of <85% should be avoided as it may lead to nonunion development.

UNDERSTANDING TECHNIQUES OF AND ATTITUDES TOWARDS PREOPERATIVE RISK ASSESSMENT

Purpose of study In the era of big electronic health data, formal surgical risk assessment tools have been developed to predict risk of adverse postoperative outcomes. Patients desire to understand their risks and report improved shared decision-making with use of such tools. However, these risk assessment tools are underutilized. We surveyed surgeons’ techniques of and attitudes towards preoperative risk assessment.
VASCULAR ANOMALIES, THROMBOEMBOLISM, AND DIETARY SUPPLEMENT USE AMONG COLLEGE STUDENTS AND ATHLETES: A LITERATURE REVIEW

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Purpose of study Recent reports have suggested that 15–30% of dietary supplements do not list the ingredients or the amounts posing a risk of adverse health events to the users. Therefore, gaining understanding about the patterns of dietary supplement use among youth is very important. The purpose of this study was to investigate the characteristics associated with supplement use among college students and athletes.

Methods used A literature review using Pubmed and Google Scholar with the keywords ‘Use of Dietary Supplements among College Student’, ‘Dietary Supplements College Athletes’, ‘College Student Supplement Use’ was conducted. Only studies published after 2003 that incorporated questionnaires that categorized the frequency of supplement use in otherwise healthy college students were included in our analysis.

Summary of results Of the 17 articles, only 10 satisfied our inclusion criteria with 6 studies involving college students and 4 involving college athletes exclusively. Prevalence of supplement use ranged from 26.3%-86.9% among college students and 76.8%-89.0% among college athletes. The most dietary supplement used were energy or caffeine containing drinks (48-80%), vitamins/mineral (18-80%) and protein supplements (17-42%). The reasons given for use of supplements were: to promote and maintain good health (14-73%), increase energy (29-50%), and improve strength and muscle (20-43%). The most popular source of information or influence for supplements among women was their families and for men was the internet. One limitation noted among the studies was the wide variation in the frequencies mainly due to the differences in reporting of short-term vs. long-term or recurrent use.

Conclusions Supplement use is prevalent among college students and college athletes. With the increasing availability and access to these supplements, it is critical that healthcare professionals are aware of the patterns of usage and educate patients about potential risks.
professionals inquire about their use and serve as an important resource to provide education and awareness. Students should be educated to consume healthy diets consistent with accepted nutritional requirements and discouraged from using supplements to substitute for other healthy nutrition or behaviors.

Purpose of study Scoliosis often requires the wearing of a back brace. There are a variety of braces that are used, and there is no data one brace is superior in terms of efficacy. The quality of life is negatively affected by the wearing of a scoliosis brace but little data is available on how different types of scoliosis braces affect the quality of life. We hypothesized that the type of scoliosis brace used in treatment impacted the quality of life of patients with scoliosis.

Methods used An online survey was conducted to gather data on patients with scoliosis; all participants granted informed consent. Collected data included demographic data, clinical information, treatments administered, compliance, patient experience including friendships, teasing, attitude towards scoliosis back brace, and involvement in choice of the brace used in their treatment. In addition, the Scoliosis Research Society-24 (SRS-24) instrument was used to determine subject satisfaction and performance. The SRS-24 instrument is a validated instrument used in adolescents with idiopathic scoliosis to determine satisfaction and performance after treatment. The SRS-24 measures pain, self-image, function, level of activity, and satisfaction.

Summary of results Data collection is ongoing; to date 32 scoliosis patients have responded. Almost all respondents (96.9%) wore a brace as part of their treatment, 43.8% of patients had surgery, and 38.7% were prescribed exercises as part of their treatment. Among respondents, 59% wore a Boston brace, 19% Rigo Cheneau, 13% Providence, 3% Charleston Brace, and 6% did not know their type of brace. Doctors chose the brace used in treatment for most patients (77%); parent(s) or guardian(s) chose the brace in 19.4% of patients. Satisfaction with wearing a scoliosis brace was low with 28.1% of respondents reporting they ‘hate’ their brace, and 43.8% stating they ‘dislike’ their brace. The total mean SRS-24 score was 3.6. We did not find a correlation between type of brace and SRS-24 scores.

Conclusions Scoliosis and the use of a back brace has an impact in the quality of life of adolescent females. Despite this most girls do not have a say in the choice of their scoliosis back brace. There was no correlation between the type of brace worn and attitudes towards wearing the brace on the SRS-24 score.

Purpose of study The purpose of this study was to identify the demographic characteristics, analyze obesity related comorbidities, and identify weight outcomes of patients seen in the UCLA Fit for Healthy Weight Program (Fit Clinic), a multidisciplinary pediatric weight management clinic.

Methods used A retrospective chart review was conducted of 153 pediatric patients in the Fit Clinic between August 2014 and September 2016. The mean changes in BMI z-scores over time was evaluated using a mixed effects linear regression model allowing for random intercepts and slopes.

Summary of results The average baseline BMI was 31.94 kg/m² ± 7.97. In the sample, 11% (n=17) were overweight and 89% (n=136) were obese, including 56% (n=85) who were morbidly obese. The patients had an average of 4 to 5 problem diagnoses per visit. The prevalence of comorbidities was higher in the older age group (figure 1). The most common comorbidities for both older and younger age groups included dyslipidemia (50.0% vs. 45.3%), prediabetes (35.9% vs. 21.3%), and obstructive sleep apnea (10.3% vs. 4.0%). The mean BMI z-score tended to decrease over time, mean slope = -0.05 over 6 months (p=0.05).

Conclusions Many Fit Clinic patients manifest obesity-related comorbidities and are at risk of adverse health outcomes related to cardiometabolic morbidity. The number of comorbidities increased with age, highlighting that early treatment may be imperative in preventing obesity-related comorbidities in older adolescents and adults. Fit Clinic patients’ overall mean BMI z-score tended to decrease over time, demonstrating a promising effect of a multidisciplinary approach to managing pediatric obesity.
IMPLEMENT Asthma Collaborative study based out of The University of California, San Francisco.

Methods used Retrospective chart reviews of randomized ambulatory clinic pediatric patients with a diagnosis of asthma or reactive airway disease before interventions and after interventions will be used to compare the rates of personalized electronic medical record (EMR)-based Asthma Action Plan provision. This will assess the efficacy of our interventions, which focus on resident-directed education. A portion of the data is included in the IMPLEMENT Collaborative study, however a larger population of patients is included in our dataset. We will also administer pre- and post-surveys to assess for resident awareness and comfort level with outpatient asthma management and to assess for potential areas of ongoing improvement.

Summary of results We have seen an overall increase in the provision of personalized Asthma Action Plans compared to prior to the implementation of an EMR-based system. This study began in March 2018. Asthma Action Plans were provided at an average rate of approximately 20% as of June 2018; this has increased to approximately 35% as of August 2019.

Conclusions Continuing resident physician education and implementation of a standardized EMR-based system for provision of personalized written Asthma Action Plans has led to an increased rate of utilization of this intervention. With further interventions, we expect to see further increase in the frequency of provision of a personalized Asthma Action Plan.

Abstract 233 Figure 1 Patient presented with bilateral helical rim deformity. First image is the left ear prior to treatment at 2 weeks old. Second image is the splint applied. Third image is the final result after two months of treatment.

CORRECTION OF NEONATAL AURICULAR DEFORMITIES WITH DUODERM®: A SIMPLE TECHNIQUE

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Case report Ear molding in neonates has been shown to successfully correct congenital auricular deformities. There are several currently available molding techniques. However, commercially available molding devices (e.g. EarWell and Ear Buddy) can be costly, and their alternatives have limited customizability. We selected illustrative cases seen in the BC Children’s Hospital plastic surgery clinic in the past 5 years to present the outcomes of a molding technique using cost-effective and customizable materials for common deformations (Stahl’s ear, constricted ear, and prominent ear). DuoDERM® Extra-thin, Steri-strips®, and 3M™ Kind Removal Silicone tape are used to splint the ear in a preferred position. The DuoDERM® is rolled into a putty, placed in the ear, and secured with tapes. This treatment is initiated in the clinic, with weekly splint changes carried out at home by caregivers, and intermittent follow-up appointments. Primary physicians/pediatricians can start this treatment as early as possible in the neonatal period, potentially reducing the need for future surgery. DuoDERM® molding is a safe, inexpensive, highly customizable, and simple way to correct auricular deformities.

Cardiovascular III
 Concurrent session
 8:00 AM
 Friday, January 24, 2020

234 A RARE PRESENTATION OF EOSINOPHILIC THROMBOTIC MYOCARDITIS

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Purpose of study A 62-year-old very healthy gentleman with a history of seasonal allergies presented to the hospital with stroke symptoms. Initial complete blood count showed an eosinophilic count of 44 thousand. The patient’s transesophageal echocardiogram showed a left atrial appendage thrombus. For initial concern of cardiac embolic phenomenon, anticoagulation therapy was started but a subarachnoid hemorrhage soon developed. His course complicated quickly to include renal failure. The patient was transferred to a tertiary center where he was dialyzed and improved significantly. Despite improvements and steroid treatment, he remained in critical condition.

Methods used Patient chart and literature review of ‘Eosinophilic Myocarditis with Thrombosis’. Summary of results His cardiac magnetic resonance imaging (MRI) and biopsy both suggested confirmatory diagnosis of eosinophilic myocarditis with left ventricular ejection fraction of 35% with acute congestive heart failure symptoms. He received a dose of cyclophosphamide and heart failure medications. Eosinophilic myocarditis with thrombotic phenomena is a very rare condition and presents as a stroke phenomenon and often found to have a left atrial appendage clot. There are 3 stages of eosinophilic myocarditis progression: inflammatory, thrombotic and fibrotic.

Conclusions The patient is doing very well on cyclophosphamide to date.
**Purpose of study** It is difficult to evaluate diastolic dysfunction on cardiac MR since to date this has required the performance of phase contrast imaging across the mitral valve to generate data similar to echocardiography. The purpose of this study is to determine if the myocardial bounce seen during atrial contraction at end diastole corresponds to diastolic and systolic function, and if loss of the bounce could be predictive of both diastolic and systolic heart failure.

**Methods used** 146 consecutive cardiac MR exams between Sept. and Dec. 2017 were selected for analysis. The bounce was graded by two blinded observers, and the change in LV diameter pre and post bounce was measured. The bounce was defined as the rapid change in LV volume that occurs at the end of diastole during atrial contraction just prior to systolic ejection. Inter-reader agreement was summarized using Cohen’s kappa. Spearman’s rank correlation coefficient was used to evaluate associations between bounce grade and cardiac physiology parameters.

**Summary of results** Overall agreement was good with unweighted kappa = 0.69 (95% CI: 0.60–0.79). Bounce grade was significantly correlated with the average change in LV diameter before and after the bounce (Spearman’s rho = 0.76, p<0.001). Median diameter changes were 0.0, 1.9, and 4.2 mm in grades 0 (no bounce), 1 (small bounce), and 2 (normal), respectively. Bounce grade was significantly correlated with LV EF (Spearman’s rho = 0.43, p<0.001). Median EF was 44%, 51%, and 58% in grades 0, 1, and 2, respectively. Of the 88–89 patients who had E/A ratio or E/e’ ratio measured, bounce grade was also significantly correlated with E/A ratio (r = -0.24, p=0.034) and E/e’ ratio (r = -0.24, p=0.022), with lower grades having higher ratio values on average.

**Conclusions** The simple observation of a myocardial bounce during cine loop review of cardiac MR exams was predictive of diastolic and systolic cardiac function. Lack of myocardial bounce was highly associated with both systolic and diastolic dysfunction. The subpopulation of patients with loss of myocardial bounce and normal EF appear to represent patients with early diastolic dysfunction, but further evaluation will be needed with more patients with known diastolic heart failure.
defibrillator placement, and stroke), and 1-year freedom from rejection, including any treated rejection (ATR), acute cellular rejection (ACR), and antibody-mediated rejection (AMR).

Summary of results Patients with severe PGD had decreased 3-year survival, 1-year freedom from any treated rejection, and 3-year freedom from NF-MACE compared to those patients who did not have severe PGD. There were no significant differences between the two groups in terms of 3-year freedom from CAV and freedom from DSA.

Conclusions Severe PGD appears to have increased mortality and morbidity with more rejection and more NF-MACE. More intense therapies to offset the inflammatory response from severe PGD should be investigated.

Summary of results Patients in the lowest quintile of LDL cholesterol did not appear to have significantly improved outcomes compared to the other quintiles.

Conclusions Lower levels of LDL cholesterol at 1-year post-transplant do not appear to have beneficial outcomes although all patients were on statin therapy. Aggressively lowering cholesterol levels may not be indicated in this patient population. Larger studies are warranted to confirm these results.

Purpose of study Low-density lipoprotein (LDL) cholesterol has been demonstrated to adversely affect patients with underlying coronary artery disease. In heart transplantation (HTx), LDL cholesterol rises due to the use of corticosteroids and calcineurin inhibitors. From two randomized trials in HTx, it has been demonstrated that statins can lower LDL but also have an immunomodulatory effect by decreasing severe rejection and cardiac allograft vasculopathy (CAV) thus improving survival. It is not been established whether lowering LDL cholesterol to <70 mg/dL can result in improved HTx outcomes.

Methods used Between 2010–2016, we assessed 109 HTx patients and assessed their LDL cholesterol at 1-year post-transplant. All patients were on statin therapy and patients were divided into quintiles based on their cholesterol levels. The groups were then compared for subsequent 3-year outcomes, including survival, freedom from CAV (as defined by stenosis >50%), freedom from non-fatal major adverse cardiac events (NF-MACE, defined as myocardial infarction, percutaneous coronary intervention/angioplasty, new congestive heart failure, pacemaker/implantable cardioverter-defibrillator placement, and stroke), and 1-year subsequent rejection including any treated rejection (ATR), acute cellular rejection (ACR), and antibody-mediated rejection (AMR). PCSK9 was not used in this study.

Conclusions Severe PGD appears to have increased mortality and morbidity with more rejection and more NF-MACE. Larger studies are warranted to confirm these results.
Abstract 239 Table 1

<table>
<thead>
<tr>
<th>Endpoints</th>
<th>Group A (Control):</th>
<th>Group B:</th>
<th>Group C:</th>
<th>Group D:</th>
</tr>
</thead>
<tbody>
<tr>
<td>(1-Year)</td>
<td>PRA 0% (n=158)</td>
<td>PRA 1-25% (n=55)</td>
<td>PRA 26-50% (n=57)</td>
<td>PRA &gt;50% (n=100)</td>
</tr>
<tr>
<td>Survival</td>
<td>90.5%</td>
<td>85.5%</td>
<td>91.2%</td>
<td>94%</td>
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<tr>
<td>Freedom from CAV</td>
<td>94.9%</td>
<td>98.2%</td>
<td>93%</td>
<td>94%</td>
</tr>
<tr>
<td>Freedom from NF-MACE</td>
<td>82.9%</td>
<td>92.7%</td>
<td>86%</td>
<td>86%</td>
</tr>
<tr>
<td>Freedom from ATR</td>
<td>83.5%</td>
<td>89.1%</td>
<td>89.5%</td>
<td>81%</td>
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<tr>
<td>Freedom from ACR</td>
<td>93.7%</td>
<td>92.7%</td>
<td>96.5%</td>
<td>93%</td>
</tr>
<tr>
<td>Freedom from AMR</td>
<td>93.7%</td>
<td>94.5%</td>
<td>93%</td>
<td>85%*</td>
</tr>
<tr>
<td>Freedom from DSA</td>
<td>88.6%</td>
<td>83.6%</td>
<td>75.4%**</td>
<td>50.0%***</td>
</tr>
</tbody>
</table>

*Significant with p=0.020 when compared to Control  **Significant with p=0.024 when compared to Control  ***Significant with p=0.001 when compared to Control

Purpose of study Mechanical circulatory support (MCS) is increasing for patients with advanced heart disease. A common complication is a driveline infection. It is also known that approximately 30% of MCS patients develop circulating antibodies (known as sensitization) due to blood transfusions. Becoming sensitized narrows the donor pool due to incompatible donors. For these patients, desensitization therapy is important to allow transplantation to occur with a compatible donor. There is concern that the use of desensitization therapy might worsen infection in MCS patients with an existing driveline infection.

Methods used Between 2007 and 2018, we assessed 45 MCS patients who developed a driveline infection. These patients were divided into those who were sensitized (PRA ≥ 10%, n=22) and those who were not sensitized (n=23). The sensitized patients were further divided into those that received desensitization therapy (n=7) and those that did not (n=15). The groups were monitored for the ensuing 1 year after the driveline infection for survival, freedom from readmission, freedom from new driveline infection, and freedom from any infection.

Summary of results There was no significant difference between sensitized and non-sensitized MCS patients with a driveline infection in terms of 1-year survival, freedom from hospital readmissions, and freedom from any infection subsequent to the initial driveline infection. Additionally, when comparing the subgroup of sensitized patients who received desensitization therapy and those that did not, there was no difference in outcome.

Conclusions Desensitization appears to be safe in MCS patients with an existing driveline infection to potentially expand the compatible donor pool.

Abstract 240 Table 1

<table>
<thead>
<tr>
<th>Endpoints (at 1-Year)</th>
<th>Driveline Infection, PRA ≥10% (n=22)</th>
<th>Driveline Infection, Non-sensitized (n=23)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Survival</td>
<td>95.2%</td>
<td>100.0%</td>
<td>0.317</td>
</tr>
<tr>
<td>Freedom from Readmission</td>
<td>30.4%</td>
<td>4.6%</td>
<td>0.102</td>
</tr>
<tr>
<td>Freedom from New</td>
<td>79.3%</td>
<td>75.9%</td>
<td>0.848</td>
</tr>
<tr>
<td>Driveline Infection</td>
<td>Freedom from Any Infection</td>
<td>49.5%</td>
<td>54.8%</td>
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</tbody>
</table>

Abstract 240 Table 2

<table>
<thead>
<tr>
<th>Endpoints (at 1-Year)</th>
<th>Driveline Infection, PRA ≥10%, Desensitization Therapy (n=7)</th>
<th>Driveline Infection, PRA ≥10%, No Desensitization Therapy (n=15)</th>
<th>P-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Survival</td>
<td>100.0%</td>
<td>92.9%</td>
<td>0.498</td>
</tr>
<tr>
<td>Freedom from Readmission</td>
<td>14.3%</td>
<td>0.0%</td>
<td>0.328</td>
</tr>
<tr>
<td>Freedom from New</td>
<td>83.3%</td>
<td>77.9%</td>
<td>0.727</td>
</tr>
<tr>
<td>Driveline Infection</td>
<td>Freedom from Any Infection</td>
<td>33.3%</td>
<td>71.3%</td>
</tr>
</tbody>
</table>

Purpose of study Mechanical circulatory support (MCS) is increasing for patients with advanced heart disease. A common complication is a driveline infection. It is also known that approximately 30% of MCS patients develop circulating antibodies (known as sensitization) due to blood transfusions. Becoming sensitized narrows the donor pool due to incompatible donors. For these patients, desensitization therapy is important to allow transplantation to occur with a compatible donor. There is concern that the use of desensitization therapy might worsen infection in MCS patients with an existing driveline infection.

Methods used Between 2007 and 2018, we assessed 45 MCS patients who developed a driveline infection. These patients were divided into those who were sensitized (PRA ≥ 10%, n=22) and those who were not sensitized (n=23). The sensitized patients were further divided into those that received desensitization therapy (n=7) and those that did not (n=15). The groups were monitored for the ensuing 1 year after the driveline infection for survival, freedom from readmission, freedom from new driveline infection, and freedom from any infection.

Summary of results There was no significant difference between sensitized and non-sensitized MCS patients with a driveline infection in terms of 1-year survival, freedom from hospital readmissions, and freedom from any infection subsequent to the initial driveline infection. Additionally, when comparing the subgroup of sensitized patients who received desensitization therapy and those that did not, there was no difference in outcome.

Conclusions Desensitization appears to be safe in MCS patients with an existing driveline infection to potentially expand the compatible donor pool.

Abstract 241

Effectiveness of Intravenous Immunoglobulin versus Infliximab in Patients with Refractory Kawasaki Disease

Purpose of study It is unclear whether the use of Infliximab (IFX) or a second dose of intravenous immunoglobulin (IVIG) is more effective in treating patients with IVIG-resistant Kawasaki Disease (KD). The purpose of this study is to investigate whether retreatment with IFX or IVIG is associated with a greater and more rapid resolution of fever and fewer adverse coronary outcomes in patients with refractory KD.

Methods used A literature review using Pubmed and Google Scholar with the keywords Infliximab, IVIG-resistant Kawasaki
Disease, and Refractory Kawasaki Disease was conducted. Only studies that directly compared retreatment with IVIG versus IFX were used. Of these studies, only those that evaluated coronary outcomes, response rates, and febrile periods for each retreatment were used.

Summary of results Of the 16 articles that we found, only 4 satisfied our inclusion criteria (see table 1 below). Our study was limited due to most studies focusing on treatment for patients with non-refractory KD. Additionally, many studies combined different treatment options together and were thus excluded from this study. All studies that tested IFX found a correlation between Infliximab administration and greater resolution of fever along with shorter febrile periods. The correlation between coronary abnormalities and the administration of IFX or IVIG remains inconclusive.

Conclusions Our review suggests that IFX could be more successful in alleviating fevers versus retreatment with IVIG in refractory KD patients. The occurrence of coronary abnormalities and adverse events in retreatment with both drugs was not significant. Further research with larger sample sizes and longer follow up periods must be conducted to fully confirm these results.

Case reports I Concurrent session 8:00 AM Friday, January 24, 2020

242 COLON CANCER PRESENTING AS A SPLENIC ABSCESS WITH ASSOCIATED FUSOBACTERIUM MORTIFERUM BACTEREMIA

E Fitzsimmons, T Tonica, C Donath, S Ragland, R University School of Medicine, Bridgetown, Barbados and, Kem Medical Center, Bakersfield, CA.

Abstract 241 Table 1  Efficacy of retreatment with IVIG vs. IFX in refractory kawasaki disease

<table>
<thead>
<tr>
<th>Study</th>
<th>Location</th>
<th># of Patients Studied</th>
<th>Median febrile period (in days)</th>
<th># of patients who became afebrile</th>
<th>Response Rate</th>
<th># of patients with adverse coronary outcomes</th>
<th>Median echocardiogram z-scores</th>
<th>Echocardiogram z-score p-value</th>
<th>Follow-up length</th>
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<tbody>
<tr>
<td>IVIG</td>
<td>IFX</td>
<td>IVIG</td>
<td>IFX</td>
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<td>IFX</td>
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<tr>
<td>Burns 2008 US 24 NR NR 8/12 (67%) 11/12 (92%) p&lt;0.05 1/12 4/12 2.2 3.5 NS Study entry, 1–2 &amp; 6–8 weeks</td>
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<tr>
<td>Son 2011 US 106 10 8 65.06 17/20 (76%) (85%) 10.20 (17%) (24%) p=0.030 2/86 7/20 1.60 1.76 NS Study entry, 1–10 weeks</td>
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<tr>
<td>Yoon 2016 Korea 43 10 8 22.93 14/15 (65.3%) (83%) 10.20 (17%) (24%) p=0.042 4/33 1/15 NR NR NR</td>
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<tr>
<td>Mori 2018 Japan 31 9.3 7.7 6/15 12/16 (40%) (75%) 2.5 1.9 NS Study entry &amp; 3, 7, 14, 21, and 56 days</td>
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</table>

*febrile period following initial IVIG treatment until resolution of fever after retreatment; NS: Not Significant; NR: Not Reported

Case report Streptococcus bovis bacteremia is a well-known classic association for an underlying colorectal carcinoma. However, more rarely, Gram-negative bacteremia’s can also be associated with colorectal cancer. This case report documents an underlying colorectal carcinoma with a previously undescribed Fusobacterium mortiferum bacteremia. In addition to the bacteremia, this patient presented with a splenic abscess, whose formation generally requires seeding from another primary source of infection. Cultures of the splenic abscess in this case did not grow Fusobacterium mortiferum but did grow Escherichia coli and Streptococcus group C beta. Surgical intervention later confirmed suspicion of direct contact of the distal transverse colon to the spleen due to an expanding colorectal carcinoma as no other primary sources of infection were found. To the best of our knowledge, there are no documented cases of Fusobacterium mortiferum bacteremia’s associated with colorectal carcinoma in the literature. Additionally, the formation of a splenic abscess by direct contact, rather than by seeding, remains incredibly rare.
A RARE CASE OF COCCIDIOIDAL OTOMYCOSIS

HK Sandhu*, G Petersen, A Heidari. Kern Medical, Bakersfield, CA

Purpose of study Coccidioidomycosis (coccii) Immitis is found abundantly in soil in the Central Valley of California, southern Arizona, west Texas, and southern New Mexico. Around 40% of infections due to coccidioides are asymptomatic with vast majority being self-limited pulmonary infection. 1–2% of cases show dissemination beyond the lungs, most commonly to bones, lymph nodes, joints, and meninges. We here present the only case of coccidioidomycosis dissemination to the mastoid process that we experienced in our data base of over three thousand cases. Methods used Retropective case study. Summary of results A 22 year old Hispanic male with uncontrolled Diabetes Mellitus type I was diagnosed with pulmonary coccidioidomycosis prior. Coccii complement fixing titer was 1:32. He was placed on fluconazole 800 mg daily. One year later he experienced gradual onset left ear pain with radiation to left jaw and left eye and purulent drainage. He was diagnosed with left otitis externa and was treated medically. Subsequently, he presented with left sided progressive hearing loss and diffuse pounding headache, nausea and vomiting of 1 month duration. On otoscopic examination, tympanic membrane was erythematous, bulging with middle ear effusion. 512 tuning fork conclusive of conductive hearing loss on the left side. CT of the head showed complete opacification of mastoid air cell system on the left side with fluid in the middle ear. MRI showed left mastoiditis with an extradural collection. He was admitted and started on I.V. antibiotics. Upon ENT consultation operative management was conducted for left mastoiditis and otitis media with left myringotomy with insertion of tympanostomy tube. Patient remained symptomatic 10 days postoperatively therefore, left mastoidectomy and tympanoplasty was performed for chronic mastoiditis. Mastoid tissue biopsy on culture grew Cocciidomyces Immitis after 28 days. The tympanic membrane healed well and there was a gradual increase of hearing in the left ear. Patient subsequently completed 12 week course of Liposomal amphotericin B and his treatment was changed to oral posaconazole he improved and followed up in the outpatient setting. Conclusions Clinicians should be aware that coccidioidomycosis could hematogenously disseminate to any foci of body particularly in the immunocompromised host.

A CASE OF ACUTE LIVER INJURY IN THE SETTING OF ANTI-TUBERCULOSIS THERAPY FOR PRESUMPTIVE OCULAR TUBERCULOSIS

J Bhandohal*, J Bhaka, R Janiwal, R Gupta, J West, S Allday, C Anderson, B Oftedal. Kern Medical, Bakersfield, CA

Purpose of study Mycobacterium tuberculosis (MTB) is an obligate aerobic bacteria, and is the causative agent in ocular tuberculosis (TB); the oxygen tension of the choroid of the eye is one of the highest oxygen tensions in the body and is attractive for the bacillus to be seeded via hemogenous spread. Ocular TB is more common in the uveal tract, retina, and optic nerve respectively. The diagnosis is essentially clinical as the pulmonary disease is missing. Summary of results A 31-year-old Hispanic female who previously worked in a hospital in Mexico referred from her ophthalmologist with the diagnosis of ocular TB. The patient had been suffering for about 1 year with bilateral panuveitis with bilateral granulomatous infiltration. Her Quantiferon-Gold TB test, was positive with negative chest x-ray. She was initiated with rifampin, isoniazid, pyrazinamide and moxifloxacin. After about 4 months of therapy with clinical improvement she was found to have epigastric pain, nausea, vomiting, and non-bloody diarrhea; her AST/ALT were 1,424/1,584 U/L, and Bili T of 4.4 mg/dl. All anti-TB medications were stopped, and the patient was admitted. Her liver injury continued to worsen to levels virus (HIV) and ocular syphilis is an uncommon initial presentation of the infection. Methods used Retrospective case report. Summary of results A 58-year-old Hispanic male with a history significant for HIV presented to his ophthalmologist with complaints of seeing floaters in the previous three months. The floaters were associated with vision changes but no complete loss of vision or pain. Due to the patient’s HIV status and the recent vision changes, laboratory studies were performed to assess for a variety of diagnoses. The initial diagnosis of syphilis was confirmed and the patient was sent to the hospital for further management and treatment. On admission, physical exam demonstrated extraocular movements intact, pupils equal and reactive to light and accommodation with no reduction in peripheral vision. Cranial nerves II-XII were tested and intact and sensation to light touch was intact throughout. A lumbar puncture with differential and VDRL along with a serum qualitative syphilis antibody were obtained. The cerebral spinal fluid returned positive for syphilis and the patient was given a PICC line and was treated for 14 days with 4 million units of Penicillin G IV. Conclusions This case demonstrates an uncommon initial manifestation of syphilis that requires high suspicion for diagnosis and further workup to determine how invasive the infection for proper treatment protocol. Syphilis in general is more commonly seen in patients diagnosed with HIV and therefore any patient with HIV presenting with vision changes should undergo initial syphilis diagnostic testing as a delay in treatment may cause a worsening of symptoms. Finally, if ocular syphilis is the initial symptom at diagnosis, a CSF analysis should be done to determine if the patient also has neurosyphilis as this needs more aggressive treatment.
as high as 19/1957/2363 (Bili/AST/ALT). MELD as high as 23 and INR as high as 2.

She underwent liver biopsy complicated with intra-abdominal hemorrhage and embolization. Pathology showed perivenular necrosis, ballooning degeneration of hepatocytes, and mild bridging fibrosis, consistent with drug induced necrosis. Her ANA came back 1:160 suggestive of underlying autoimmune process as well. She was transferred to liver transplant center but fortunately, spontaneously start to improve with slow drop in liver injury markers. Plan is to restart her treatment only with 2 least hepatotoxic medications to finish her course of treatment.

Conclusion Clinicians should be aware of the potential acute and severe liver injury when treating patients with ant-tuberculosis medications. Due to nature of clinical diagnosis and lacking microbiological data to deescalate treatment the risk of toxicity is higher.

246 RECURRENT HOSPITALIZATION FOR CANNABINO3 HYPEREMESIS SYNDROME IN AN ADOLESCENT
K De Leon*, J Pham, T1 Chimino, Loma Linda University, Loma Linda, CA
10.1136/jim-2019-WMRC.246

Case report A 16 year-old male with history of major depressive disorder and attention deficit hyperactivity disorder presented to an outside ED with four days of recurrent episodes of nausea and vomiting associated with weight loss and decreased appetite. He was transferred due to chest pain and shortness of breath attributed to pneumomediastinum identified on chest radiograph. The patient reported at least 2 years of daily use of ‘wax’ marijuana which he smoked to maintain a constant ‘high’.

On admission the patient continued with nausea and non-bilious, non-bloody vomiting. Vital signs were reassuring with normal oxygen saturation. Physical exam was notable for crepitus over his upper left anterior chest wall but normal breath sounds and work of breathing. The abdomen was soft and non-distended but generally tender to palpation. Lab values were normal except for white blood cell count of 23.5 bil/L and slightly low sodium and chloride. Chest x-ray showed mediastial air and CT scan of the chest, abdomen, and pelvis confirmed pneumomediastinum with subcutaneous air, and ruled out intestinal obstruction. His symptoms were unresponsive to typical antiemetics such as ondansetron and promethazine, but he reported significant relief with hot water bathing. Based on his chronic cannabis use and symptoms he was diagnosed with CHS. Due to persistent symptoms, he was started on haloperidol after which he experienced relief. He was discharged home, but would go on to have multiple further encounters.

Over the course of approximately 18 months he presented eight times to various EDs reporting the same symptoms. Five of these eight ED visits resulted in hospital admission for a total of 23 hospital days. Recurring clinical findings were: pneumomediastinum, leukocytosis (as high as 30.5 bil/L), hypertension, hypokalemia, and hypochloremia. He was treated with ondansetron and promethazine with negligible benefit. He did not tolerate a trial of topica capsaicin. He was given haloperidol on three visits with reported improvement each time. He underwent a total of three abdominal CT scans, two ultrasounds, and numerous x-rays. He received intravenous antibiotics on two occasions. Upon phone interview with the patient’s legal guardian, he has been symptom free for 8 months since he quit using cannabis.

247 ONCE BUT NOT TWICE: RIPE BEFORE SNIP
VF Civelli*, M Valdez, V Narang, A Heidari, R Johnson. Kern Medical, Bakersfield, CA
10.1136/jim-2019-WMRC.247

Purpose of study Orchiectomy in the presence of unilateral mass is common practice in Urology; however histopathology post-resection may be consistent with infectious etiology, such as tuberculosis (TB) or endemic fungal sources. Here we describe a case of presumptive TB of the testicles in a patient who underwent unilateral right orchiectomy followed by spread of symptoms to the contralateral testicle post-procedure.

Methods used Retrospective single-patient case report.

Summary of results The patient is a 41-year-old Hispanic male who underwent right-orchiectomy, to rule out malignancy, at an outside facility two years prior after he presented with right sided testicular enlargement. He subsequently presented to our facility with new left testicular pain and swelling, which started approximately 6–10 months after the right orchiectomy. Examination revealed a chronic wound with exudative material at the base of an erythematous and edematous scrotum. A firm mass and scrotal tenderness were appreciated. A course of doxycycline was completed due to concern for epididymitis, which provided no relief. The pathology report from the right orchiectomy was obtained and revealed necrotizing granulomatous inflammation with a rare, questionable AFB-stained organism. Additional work-up for coccidioidomycosis, granulomatosis with polyangiitis, sarcoidosis, histoplasmosis, and brucellosis were negative. Urine AFB stains and cultures were also negative. Regardless, empiric therapy with the standard 4-drug anti-tuberculosis medication regimen (RIFPE), including rifampin, isoniazid, pyrazinamide, and ethambutol was initiated. The patient completed a 6 month course of RIFPE therapy and experienced complete resolution of all symptoms, including left testicular pain, swelling, and erythema.

Conclusions Clinicians should be mindful to include infectious etiologies, specifically TB, in the differential diagnosis for orchitis, epididymitis, and solid testicular masses, even in the setting of negative work-up. Although the urology literature describes that orchiectomies occur per standard guidelines for malignancy, in the case of spread of symptoms to the remaining testicle a comprehensive work-up for infectious etiologies and perhaps empiric therapy should be considered prior to resection.

248 A CASE OF HEPATITIS B IMMUNE ESCAPE VARIANT: IS MY SPOUSE SAFE?
A Shahr*, A Heidari, V Ball, M Valdez. Kern Medical-UCLA, Bakersfield, CA
10.1136/jim-2019-WMRC.248

Purpose of study Immune escape variance in hepatitis B is a rare phenomenon seen particularly in the Asian population. Antigenic escape occurs when the immune system is unable to respond properly to an infectious agent. In the case of Hepatitis B (HBV), this presents as a possibility when the patient has the coexistence of HBsAg and HBsAb. The risk
for malignancy, transmission rate, and other clinical significance of immune escape variance in hepatitis B is not well studied.

Methods used Retrospective Study

Summary of results 50 year old Phillipino male with a diagnosis of previously treated hepatitis B presented to our clinic for further evaluation after he had a positive hepatitis screening test. He reports no clinical symptoms including weight loss, abdominal pain, jaundice, pruritus. His laboratory result showed hepatitis B viral load of 36, HBsAg reactive, HBsAb 590, reactive HBeAg, nonreactive HBeAb and HBcAb without any clinical symptoms and had elevated liver enzymes (ALT 68, AST 39). Given the contradictory nature of the results, a repeat lab test was requested from the patient, which again showed HBeAg and HBeAb non-reactive, HBsAg reactive, HBsAb 706, Hep B DNA 73 and reactive HBcAb. Liver ultrasound showed coarse echogenicity of the liver suggest of cirrhotic liver. The coexistence of HBsAg and HBcAb indicates possible immune escape variants. Additional Hepatitis B genotype study was ordered, however was not able to be performed because the patient had a viral load <600. Since patient showed evidence of cirrhosis by ultrasound based on AASLD guideline he was started on Entecavir 1 mg daily and continued with unknown duration. After being on the medication for 6 months, repeat serology showed HBsAg reactive, HBsAb 726, Hep B DNA not detected.

Conclusions A diagnosis of immune escape variant of hepatitis B should be considered when the patient has serological findings of both the surface antigen and antibody. The family screening should be considered and immunization should be done in non-immunized at risk. However, risk of transmission of this variant even to individuals with adequate immunity is unknown and precautions are still warranted.

A CASE OF ESCHERICHIA COLI VULNERIS NECROTIZING FASCIITS

1,2A Hall*, 1,3K Choi, 1A Govindarajan, 1A Heldal. 1Ross University School Of Medicine, Miramar, FL; 2Kem Medical – UCLA, Bakersfield, CA; 3Arizona College of Osteopathic Medicine At Midwestern University, Glendale, AZ

Purpose of study Escherichia vulneris is a gram-negative, oxidase-negative, fermentative, motile rod in the family of Enterobacteriaceae based on DNA relatedness studies and biochemical reactions. Most isolates of E. vulneris have been recovered from wounds and few case studies are available characterizing its pathogenic characteristic. We are presenting a rare case of type I polymicrobial necrotizing fasciitis with of Escherichia vulneris, streptococcus anginosus and eikenella corrodens.

Methods used Retrospective case report.

Summary of results A 26 year old African American female with schizoaffective disorder, intravenous drug abuse with methamphetamine and hepatitis C presented with right arm redness and swelling for one week, the physical exam revealed a significant area of skin erythema and induration; in addition, the central area of eschar and necrosis noted on the proximal medial aspect of the right upper extremity. CT showed scattered gas bubbles in the right arm. She underwent emergent debridement and was diagnosed with necrotizing fasciitis of right chest wall and right upper arm. Her course got complicated with pneumothorax. She was started on broad spectrum antibiotics with vancomycin and piperclillin/tazobactam. Her intraoperative cultures repeatedly grew E. vulneris, streptococcus anginosus and eikenella corrodens.

E. vulneris was only resistant to ampicillin but sensitive to ampicillin/sulbactam. She underwent multiple debridements and her antibiotic regimen was desescalated to ampicillin/sulbactam. After clinical improvement she underwent successful skin graft.

Conclusions Even though Escherichia vulneris has been reported to colonize chronic wounds in this case it was isolated from intraoperative deep tissue source and was considered pathogenic. We only found one other case report of fatal E. vulneris necrotizing fasciitis.

Endocrinology and metabolism II
Concurrent session
8:00 AM
Friday, January 24, 2020

HIGH PREVALENCE OF METABOLIC SYNDROME IN ADOLESCENTS WITH KLINEFELTER SYNDROME

S DiKlotz*, N Tartaglia, M Kelsey, KJ Nadeau, S Davis. University of Colorado, Aurora, CO

Purpose of study Klinefelter Syndrome (KS) occurs in 1:600 males and is associated with high morbidity and mortality due to cardiovascular disease and diabetes. Metabolic syndrome (MetS) describes a set of features which confer an elevated risk for cardiovascular disease and diabetes. Previous studies have found that up to 50% of men with KS have MetS, but MetS has not been studied in adolescents with KS. We compared features of MetS in adolescents with KS to controls matched for sex, age and BMI-z score.

Methods used This is a matched case-control cross-sectional study in 30 males with karyotype 47,XXY who were 10–17 years of age. Cases were matched 1:1 on age and BMI-z score with healthy male controls. Waist circumference, seated systolic blood pressure (SBP), fasting blood glucose (FBG) and lipid panel were measured. MetS was defined as meeting 3 or more of the following criteria: fasting triglycerides (TG) >100 mg/dL; HDL cholesterol <50 mg/dL if <15 years old or <45 mg/dL if ≥15; FBG ≥110 mg/dL; waist circumference ≥75th percentile for age and sex; and SBP >90th percentile for sex, age and height. The proportion of participants meeting MetS criteria was compared using a Fisher exact test. Continuous variables were compared using a paired t-test or Wilcoxon matched-pairs signed rank test.

Summary of results Participants with KS (n=50) were well matched with controls (n=50) for age (14±1.7 vs. 14±1.5 years) and BMI-z score (0.31±1.3 vs. 0.36±1.2). MetS was present in 32% of adolescents with KS compared to only 12% of controls (p=0.028), correlating to an odds ratio of
MUSCLE MITOCHONDRIAL FUNCTION IN KLINEFELTER SYNDROME

S Cung*, KJ Nadeau, D Dabelea, M Cree-Green, S Davis. University of Colorado, Aurora, CO

Purpose of study Klinefelter Syndrome (XXY) occurs in 1 in 600 males, resulting in testosterone deficiency and a high prevalence of type 2 diabetes. Testosterone deficiency in men is known to cause insulin resistance and mitochondrial dysfunction is hypothesized to mediate this relationship, however mitochondrial function has not been assessed in XXY. The aim of this cross-sectional study was to evaluate mitochondrial function in adolescents with XXY and the relationship with insulin resistance.

Methods used Twenty-seven adolescent males with XXY (age 14.7±1.8 yrs) were compared to 75 controls (age 16.7±1.3 yrs). A subset of n=13 XXY boys were receiving testosterone replacement. In-vivo muscle mitochondrial function was assessed via phosphorus MR spectroscopy (31P-MRS) of the dominant-leg soleus muscle following 90 seconds of exercise. Fasting serum was analyzed for glucose and insulin to calculate HOMA-IR. Multiple linear regression was used to compare 31P-MRS outcomes (ADP and Phosphocreatine (PCr) time constants, rate of oxidative phosphorylation (Oxphos), and maximal mitochondrial function relative to mitochondrial density [Q_max]) between groups after adjusting for age differences. Within the XXY group, linear regression was used to assess the relationship of mitochondrial function variables with HOMA-IR and testosterone treatment status.

Summary of results There were no statistically significant differences in mitochondrial function outcomes between the two groups (ADP p=0.95, Oxphos p=0.19, PCr p=0.98, Q_max p=0.56). HOMA-IR did not have a linear relationship with any of the mitochondrial variables. There were also no differences within the XXY group by testosterone treatment status (ADP p=1.00, Oxphos p=0.20, PCr p=0.23, Q_max p=0.99).

Conclusions In-vivo post-exercise muscle mitochondrial function is not impaired in adolescents with XXY compared to controls and treatment with testosterone does not significantly relate to muscle mitochondrial function in XXY. The insulin resistance associated with XXY does not appear to be mediated by muscle mitochondrial dysfunction.

INCREASED DE NOVO LIPOGENESIS IN HEPATICS

H Rahat*, K Alexander, J Reisz, Y Garcia-Rayes, A Carreau, KJ Nadeau, A D’Alessandro, M Cree-Green. Univ. of CO Anschutz, Aurora, CO

Purpose of study Polycystic ovary syndrome (PCOS) affects 6-10% of women and is associated with hepatic steatosis (HS). HS is the best predictor of future diabetes and cardiovascular disease, yet the cause of HS in PCOS is unknown, difficult to diagnose, and treatments are limited. We sought to
identify unique serum markers for HS in obese PCOS adolescents.

Methods used All participants underwent fasting hormone and metabolic measures and abdominal MRI for hepatic fat, with HS defined as hepatic fat fraction ≥5.5%. Participants were divided into 4 cohorts of obese adolescents: 1) A development cohort of treatment-naive girls with PCOS and HS (N=11) or PCOS without HS (N=9); 2) a validation cohort similar to cohort 1 (PCOS nonHS N=7, PCOS HS N=7); 3) a non-PCOS cohort with HS (N=6) and without HS (N=14); 4) a PCOS cohort treated with at least 6 months of estrogen therapy with HS (N=3) and without HS (N=7). Serum samples underwent polar metabolite liquid/liquid extraction then global metabolite profiling by ultra-HPLC-MS. Samples from cohort 1 underwent untargeted analysis to determine metabolites that distinguished HS, which were confirmed with targetted analysis. Metabolites in the targeted panel were then assessed in samples from the other 3 cohorts.

Summary of results Significant differences in the serum metabolome of cohort 1 HS vs. non-HS were noted in amino acids, lipids, glycolytic, purine and heme metabolites (p<0.05). The most notably different compounds notable amino acids (notably glutamate), long chain fatty acids, biliarverdin and sphingosine 1-phosphate (S1P) were confirmed in the targeted panel. In samples from cohort 2, hierarchical clustering analysis of the top 25 metabolites by t-test confirmed results from cohort 1 with S1P, acylcarnitines, free fatty acids. Several amino acids confirmed as specific markers with ~71% accuracy. These markers were not different by HS status in cohort 3. In cohort 4, estrogen therapy normalized previously S1P, but did not resolve the excess long chain fatty acids and glutamate in HS.

Conclusions We determined that S1P, glutamate and several long chain fatty acids are uniquely elevated in HS+PCOS and S1P was lower in girls on estrogen therapy. These metabolites may be used as biomarkers to identify HS in girls with PCOS.

Methods used We illustrate the utility of AMON and SCNIC using a dataset (16s rRNA) from the gut microbiome and blood metabolome (LC/MS) of HIV positive individuals. This validation was part of a larger study of differences in fecal microbiomes in HIV and non-HIV populations. AMON uses KEGG Orthology to generate pathway enrichment and a hypergeometric test to verify the prediction of metabolites and their origin. SCNIC uses a microbiome table to build a correlation network to and groups of microbes are clustered together and collapsed into a smaller microbiome table.

Summary of results After benchmarking SCNIC, the optimal parameter values were an R-value above 0.5 for the SMD algorithm and for a gamma value above 0.1 for the LMM algorithm. With AMON applied to the data, 40 compounds were produced by bacteria alone, 58 by the host alone and 91 by both. Together, these analyses show that AMON can be used to predict the putative origin of compounds detected in a complex metabolome and SCNIC can help understand microbial dynamics.

Conclusions The limitations of integrating metabolomics and metagenomics are stark for complex microbial communities, where there are fewer genes of known function. With these tools, researchers can build a general framework to understand the specific interactions between certain microbes and the host, at a chemical level. Ultimately the understanding of microbiome interactions can influence many aspects of human health and disease through its metabolic activities, and with an understanding we can begin to develop more targeted therapies to mitigate human disease.
EARLY ADIPOITY REBOUND IN YOUTH WITH CONGENITAL ADRENAL HYPERPLASIA IS ASSOCIATED WITH INCREASED ADIPOITY AND BONE AGE IN ADOLESCENCE

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10.1136/jim-2019-WMRC.256

Purpose of study Youth with Congenital Adrenal Hyperplasia (CAH) have an increased prevalence of obesity. Adiposity rebound (AR) is a rise in body mass index (BMI) that corresponds to an increase in number and size of adipocytes. In the U.K., CAH youth exhibited an earlier mean AR by 3 years (1.7 yr) compared to the general population (~5 yr). Among healthy, unaffected youth, an earlier age at AR is predictive of future obesity and associated with advanced bone age. Our objective was to examine associations among AR, bone age advancement, and adiposity in youth with CAH, and associations with other brain and/or behavior abnormalities.

Methods used Data from 19 CAH patients were studied from the first clinic visit (prior to 2 yr), and every 6 months thereafter, until the last clinic visit. BMI (kg/m2) was calculated from length/standing height and weight at each visit. Visual inspection (VI) and cubic polynomial (CP) methods were used to determine age at AR. By VI, AR age was located at the nadir before the second rise of BMI after 1 year of age. By CP, AR age was located at the nadir before the second rise in the cubic model. Total fat and lean mass in adolescence were measured by DEXA. Waist circumference (WC), abdominal subcutaneous adipose tissue (SAT) and visceral adipose tissue (VAT) were measured by single-slice CT or 3-Tesla MRI at the umbilicus. Bone age x-ray near the time of imaging was studied for advancement beyond chronological age.

Summary of results Average age at AR for CAH youth was 3.3 ± 0.3 yr by VI, and 3.2 ± 1.2 yr by CP (p=0.9). Age at AR (by CP alone) was negatively correlated with total fat mass (r= -0.59, p=0.01), WC (r= -0.52, p=0.05), and SAT (r= -0.58, p=0.02), but not with lean mass or VAT. Age at AR was negatively correlated with bone age advancement by VI (r= -0.63, p=0.01) and CP (r= -0.45, p=0.03).

Conclusions CAH youth at our center exhibited an earlier AR by 2 years compared to the general U.S. population, and those with earlier AR had increased adiposity in adolescence. AR and bone age advancement could be causally related in CAH or explained by other factors such as glucocorticoid treatment that merit further study.

A CASE OF THYROTOXIC PERIODIC PARALYSIS

P De Mola, G Renganathan, N Hasan*, D Aguirre. Kern Medical, Bakersfield, CA

10.1136/jim-2019-WMRC.257

Purpose of study Thyrotoxic periodic paralysis (TPP) is a rare condition caused by untreated hyperthyroid disease. Patients typically present with lower extremity paralysis and associated hypokalemia. After IV KCl administration, lower extremity strength rapidly resolves, and the patient is often discharged without the causal hyperthyroid state exposed. Incidence of TPP is reported in 2% of patients with underlying thyrotoxicosis.

Methods used Retrospective case report

Summary of results 32 y.o. Hispanic male with no significant PMH presented to ED with complaints of lower extremity paralysis x 4 hours. Patient endorsed 4 similar episodes over the past 6 years, however, disease was not worked up as the paralysis rapidly resolved with IV KCl in the ER. ROS notable for 10lbs weight loss, exophthalmos, heat intolerance, diaphoresis, tremors and anxiety. His only home medication was 40mEq of KCl daily. He denied any family members sharing similar episodes, Physical exam demonstrated sinus tachycardia and mild exophthalmos.

In the ED, K+ was 2.7 and 40 mEq of IV KCl was administered, followed by another 40 mEq given orally. After three hours, his lower extremity strength recovered completely. Repeat labs showed a rapid response to KCl with K+ rising to 5.8. The working diagnosis was hypokalemic periodic paralysis (hypoKPP), however, with no family history and symptomology consistent with a hyperthyroid picture, a thyroid panel was ordered. Labs were significant for TSH <0.008, with a free T4 of 3.24. Thyroid U/S demonstrated no masses or nodules; but the gland was diffusely enlarged with increased vascularity. Case was discussed with endocrinologist, who concurred with the diagnosis of TPP. Patient was started on methimazole and propranolol with plans to see endocrinology outpatient. Unfortunately, patient was lost to follow-up.

Conclusions TPP is characterized by sudden onset of hypokalemia and subsequent lower extremity paralysis due to thyroid hormone sensitization of Na+/K+-ATPase channels. Clues that distinguish TPP from familial hypoKPP include a lack of family history of paralytic episodes, male sex, late presentation and signs of thyrotoxicosis. Treatment consists of nonselective beta-blockers, correction of hyperthyroid state and careful potassium supplementation. It is important for physicians to make the correct diagnosis as TPP is curable once euthyroid state is achieved.
Gastroenterology
Concurrent session
8:00 AM
Friday, January 24, 2020

258 MEDICAL THERAPY AND CLINICAL OUTCOMES IN ULCERATIVE COLITIS PATIENTS IN A COMMUNITY GASTROENTEROLOGY PRACTICE

A112

Biologic treated patients (41.6%) were more likely to be corticosteroid dependent (prednisone >10 mg per day for >60 days). Six patients (5%) became corticosteroid independent after a course of prednisone. Six patients (5%) became corticosteroid independent after a course of prednisone. Six patients (5%) became corticosteroid independent after a course of prednisone. Six patients (5%) became corticosteroid independent after a course of prednisone.

Purpose of study Ulcerative colitis (UC) is an idiopathic inflammatory bowel disease characterized by inflammation of the colon and rectum. Medical therapy is recommended to prevent complications and eventual surgery. The aim of this study was to understand the clinical course, medical treatment, and clinical outcomes of UC patients to identify gaps in treatment and plan for future outcome studies.

Methods used Over a one-year period (1/2017 to 1/2018), 119 UC patients were seen in an outpatient gastroenterology clinic. A retrospective chart review was conducted for demographic information as well as surgical and treatment history, including biologic and corticosteroid use and outcomes.

Summary of results A total of 119 patients were reviewed with a mean age of 45 years and mean UC duration of 9.8 years. The majority of patients had pancolitis phenotype (62%). Ten patients (8.4%) had colectomy/j-pouch surgery at an average 75 mo. from diagnosis of UC. Extraintestinal manifestations (EIMs) occurred in 13 patients (11%). UC was treated with mesalamine (45.4%), azathioprine/mercaptopurine (30.2%), and biologic (40.3%) therapies. Thirty patients (25.2%) were treated with at least 1 course of prednisone. Six patients (5%) became corticosteroid independent (prednisone >10 mg per day for >60 days). Biologic treated patients (41.6%) were more likely to receive treatment with prednisone than patients not treated with biologics (14.1%, p=0.001). Ten patients (8.4%) were hospitalized for UC related flaring (mean LOS 8 days, range 4–20). Remission of symptoms was achieved in 88.8% of patients.

Conclusions This cohort of UC patients in a community gastroenterology practice were older with more extensive disease phenotype (pancolitis) and had disease with durations longer than expected. Extraintestinal manifestations of UC occurred in 11% of patients and were likely underreported. Forty percent of UC patients were treated with biologic therapy. Corticosteroid use was common with 1 in 4 patients receiving at least 1 course of treatment. Corticosteroid dependent patients were uncommon. Remission was achieved in 89% of patients. No conclusions can be made comparing the effectiveness of different medical therapies due to small patient sample size.

259 EFFECTIVENESS OF FECAL MICROBIOTA TRANSPLANT IN PATIENTS WITH ULCERATIVE COLITIS: A LITERATURE REVIEW

Purpose of study Research suggests that alteration in gut bacterial colonies may play a role in the development of ulcerative colitis and/or its progression. The purpose of this study was to compile the literature regarding the effectiveness of Fecal Microbiota Transplant (FMT) in the management of ulcerative colitis (UC).

Methods used A comprehensive search using keywords such as: ulcerative colitis, FMT, and microbiota transplant was performed via Pubmed, Cochrane and checking the references of published articles. Inclusion criteria included prospective studies of adult patients diagnosed with active UC with at least 10 subjects that measured clinical outcome.

<table>
<thead>
<tr>
<th>Abstract 259 Table 1</th>
<th>Effectiveness of FMT in refractory UC</th>
</tr>
</thead>
<tbody>
<tr>
<td>1st Author, Year, Country</td>
<td>Control</td>
</tr>
<tr>
<td>Costello, 2019, Australia</td>
<td>Auto-FMT (n=35)</td>
</tr>
<tr>
<td>Cui, 2015, China</td>
<td>Self-controlled (n=15)</td>
</tr>
<tr>
<td>Ishikawa, 2017, Japan</td>
<td>Antibiotic-only (n=19)</td>
</tr>
<tr>
<td>Jacob, 2017, USA</td>
<td>Self-controlled (n=20)</td>
</tr>
<tr>
<td>Moayyedi, 2015, Canada</td>
<td>Water enema (n=34)</td>
</tr>
<tr>
<td>Paramsothy, 2019, Australia</td>
<td>Placebo enema (n=43)</td>
</tr>
<tr>
<td>Rossen, 2015, Netherlands</td>
<td>Auto-FMT (n=25)</td>
</tr>
<tr>
<td>Tian, 2019, China</td>
<td>Self-controlled (n=20)</td>
</tr>
</tbody>
</table>

*Mayo score is a clinical index used to assess the severity of UC. Lower scores indicate less symptoms.
Summary of results Of 16 studies found, 8 met our inclusion criteria (table 1 below). Based on the limited data, FMT via enemas and colonoscopies demonstrated significant positive effects, whereas FMT via a nasoduodenal tube yielded insignificant improvement. The timing of data collection was very variable and ranged from 4 weeks to 18 months, and some studies lacked long follow-up periods. Common adverse effects include nausea, borborygmus and discomfort, while less than 10% of participants experienced more serious adverse effects such as intolerance to the FMT procedure and severe worsening of symptoms.

Conclusions Our review suggests a possible role in FMT in certain patients who don’t respond to conventional therapy. Limitations of this review included variation in the type and frequency of FMTs delivered as well as the timing of assessment in different studies. Future controlled studies with a standard method of FMT and longer follow-up periods are warranted.

260 PREVALENCE AND CLINICAL SIGNIFICANCE OF OCCULT HCV INFECTION

T Saito*. University of Southern California, Los Angeles, CA

10.1136/jim-2019-WMRC.260

Purpose of study HCV infection is one of the major causes of end-stage liver diseases, which account for over 15,000 deaths annually in the US. The eradication of pathogen is the only definitive medical intervention. Currently, the successful clearance of HCV is determined by the undetectable viral genome in serum 12 weeks after the completion of the antiviral therapy, referred to as sustained virological response (SVR). Recently, anti-HCV therapy has revolutionized due to the development of direct-acting antivirals (DAA). DAA therapy offers SVR in nearly all patient with negligible toxicities. In contrast, the previous standard regimen, interferon is associated with intolerable toxicity and offers SVR only 60% of patients. Given this significant improvement, the number of patients who attained SVR is exponentially increased worldwide.

It is known that HCV could maintain dormant lifecycle in the liver tissue or PBMC after achieving SVR. This condition is called ‘occult HCV infection’ (OCI). The incidence of OCI is extremely rare in patients who achieved SVR with IFN, however, the prevalence of OCI in the population who attained SVR with DAA remain unexplored. Therefore, this study aimed to delineate the prevalence and clinical significance of OCI among patients who attained SVR with DAA.

Methods used Total 50 subjects who developed abnormal liver function test, liver cancer, or liver failure after achieving SVR were enrolled. Liver tissue and PBMC were collected and subjected to the detection of HCV genome via RT-qPCR. In addition, histopathological and electron microscopic (EM) analysis of the liver tissue were performed.

Summary of results 6 subjects (12%) had detectable HCV genome in the liver tissue. HCV-RNA was also detected in PBMC of 3 subjects (6%). 2 subjects (4%) had detectable HCV genome in both liver tissue and PBMC. The histopathological analysis did not demonstrate any specific findings associated with the presence or absence of OCI. In contrast, EM analysis of OCI liver tissue showed unique features of cell injury and HCV replication.

Conclusions Our study revealed a high prevalence of OCI among patients who achieved SVR with DAA. In addition, EM analysis of the liver tissue indicated that the presence of OCI might lead to the further progression of liver disease, suggesting the potential benefit of reinitiating antiviral therapy in those with evidence of OCI.

261 EXAMINATION OF WNT SIGNALING AS A THERAPEUTIC TARGET FOR PANCREATIC DUCTAL ADENOCARCINOMA USING A PANCREATIC TUMOR ORGANOID LIBRARY

HJ Hawkins*, BW Yacob, CD Brindley, WA Messersmith, PJ Dempsey, TM Pitts. University of Colorado, Aurora, CO

10.1136/jim-2019-WMRC.261

Purpose of study Pancreatic ductal adenocarcinoma (PDAC) commonly presents at advanced stages and is refractory to most treatment modalities, making it one of the most lethal cancers. Although the low tumor cellularity and high desmoplastic response convolutes the relationship between genotype and biological phenotypes, gene mutations associated with PDACs have been identified. Wnt pathway mutations are rarely detected in PDAC, but Wnt signaling is activated by pancreatic duct ligation injury and plays a critical role in the proliferation and chemotherapeutic resistance in other cancers. Patient derived pancreatic tumor organoid libraries (Ptol) allow for more accurate investigation of the biological phenotypes that might lead to therapies that further improve survival. This study aims to subclassify PDAC organoids based on Wnt dependency to determine if combinatory treatment with Wnt inhibitors and chemotherapy would serve as a feasible treatment.

Methods used Nine PDAC organoids were grown in Human Pancreatic Stem Cell medium. Minimal media conditions required to maintain growth was assessed with deletions of various niche factors. For confirmation of Wnt inhibition, organoids grown in minimal media were treated with three Wnt inhibitors. Growth was assessed with CellTiter Glo 3D.

Summary of results Minimal media conditions, growth factor dependency, and Wnt dependency determined via Wnt inhibition were determined as described above. Five organoids demonstrated Wnt dependency. Two organoids demonstrated a Wnt independent phenotype. Minimal growth media was determined for two additional organoids, but Wnt dependency has yet to be assessed.

Conclusions Based on the results obtained, each pancreatic organoid demonstrated different niche factor dependencies providing an avenue for targeted therapy, particularly with Wnt inhibition. WES and gene expression analysis of each organoid will allow for correlation between genotype and Wnt (in)dependency observed in vivo. Combinatory treatment with Wnt inhibition and chemotherapy will be assessed in vitro and confirmed in patient-derived xenograft models.
Purpose of study: Cannabinoid (marijuana) usage has increased significantly in recent years, especially after its recent legalization. The objective of the study is to determine the association between cannabinoid use with hyperemesis or cyclic vomiting syndrome (CVS).

Methods used: A literature-based review of articles was performed through PubMed, Cochrane, and Google Scholar database using key words such as ‘Cannabinoid’ ‘Hyperemesis Syndrome’, and ‘Cyclical Vomiting Syndrome’. The studies included took into account the confounding variables or excluded patients with other illnesses that could lead to hyperemesis.

Summary of results: We found 6 studies that matched our inclusion criteria (See table 1). Duration of cannabinoid use was variable among the studies. Studies showed that cannabnoioid use has increased in recent years. In addition, cannabinoid users were more likely to be hospitalized or visit Emergency Department (ED) due to hyperemesis in comparison to non-users. Some studies noted that patients with cannabinoid associated vomiting reported compulsive hot showering to relieve vomiting. The reports suggested that the symptoms of hyperemesis stopped after discontinuation of cannabinoids.

Conclusions: Our literature review suggests that cannabinoid use is associated with hyperemesis syndrome. As the workup for diagnosis of hyperemesis syndrome could be very costly, healthcare professionals should have a high index of suspicion for cannabinoid use in patients who present with hyperemesis.

Abstract 262 Table 1: Association of cannabinoid use and hyperemesis syndrome

<table>
<thead>
<tr>
<th>First Author, Year, Location</th>
<th>Method of study/Variable Measured</th>
<th>Number of Cases vs Controls, and Age Range</th>
<th>Frequency of variable in Subjects vs Controls</th>
<th>P value (other stats)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hernandez, 2019, Canada</td>
<td>Compared use of cannabis in patients who presented to ED with hyperemesis syndrome</td>
<td>37 cannabis users vs. 336 control (18-55 yrs)</td>
<td>Hyperemesis in 16 (42.3%) of users 62 (18.5%) of non-users</td>
<td>0.001</td>
</tr>
<tr>
<td>Venkatesan, 2019, U.S.</td>
<td>Compared cannabis use in patients with CVS</td>
<td>140 patients with CVS (37±13 years)</td>
<td>41% percent of patients with CVS were current cannabis users, with 21% reported regular use</td>
<td>Not Reported</td>
</tr>
<tr>
<td>Bhandari, 2018, U.S.</td>
<td>Compared cannabis use in hospitalized patients with CVS vs non-CVS</td>
<td>4715 CVS vs 2,359,247 non-CVS Hospitalizations, all ages (mostly between 18–45 yrs)</td>
<td>Cannabis use in 13% of CVS vs 1.7% of non-CVS</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Patel, 2015, U.S.</td>
<td>Compared use of cannabis in patients hospitalized with persistent vomiting (PV) vs no-PV</td>
<td>47,753,472 no-PV vs 55,549 with PV, (15 to 54 yrs)</td>
<td>Cannabis use risk no-PV=3.9 vs PV=20.8</td>
<td>p&lt;0.001</td>
</tr>
<tr>
<td>Kim et al. 2015, U.S.</td>
<td>Comparison of CVS pre and post-liberalization of marijuana in patients seen in ED</td>
<td>41 visits for CVS Pre-liberalization vs 87 visits Post-liberalization</td>
<td>Cannabis use in 7 (17%) Pre-liberalization vs 37 (43%) Post-liberalization</td>
<td>P&lt;0.05</td>
</tr>
<tr>
<td>Habboushe, U.S. 2019</td>
<td>Survey to identify Cannaboid Hyperemesis (CH) according to cannabinoid use</td>
<td>2127 patients (18–49 year old) were surveyed for marijuana usage</td>
<td>51 (23.9%) of frequent cannabinoid users (&gt;20 days/month) met the definition of Cannaboid Hyperemesis Syndrome</td>
<td>Not Applicable</td>
</tr>
</tbody>
</table>
A LARGE COHORT OF CNKSR2-RELATED DISEASE PROVIDING INSIGHTS INTO NATURAL HISTORY AND THE MUTATIONAL SPECTRUM

Higa, Singh, Massiello, Foster, Shen. Common Regional Medical Center, Fresno, CA; CNKSR2 Family Support Group, Reedley, CA; UCSF Fresno, Fresno, CA.

Purpose of study A pathogenic variant in CNKSR2 (connector enhancer of kinase suppressor of Ras-2) was uncovered in a boy with a seizure disorder and delays especially in language. We have subsequently recruited the largest known cohort that doubles the number of affected individuals, and triples the number of suspected pathogenic variants, with the aim of improving our understanding of the neurodevelopmental and seizure characteristics of CNKSR2-related disease and its mutational spectrum.

Methods used The proband was analyzed by clinical whole exome through a commercial laboratory. Clinical and molecular information from additional cases were provided through parents active in a CNKSR2 family support group.

Summary of results CNKSR2-related disease is an X-linked disorder that results in a neurobehavioral phenotype consisting of intellectual disability and seizures. The delay in developmental progression can be severe, particularly so in language skills. There are characteristic EEG patterns during sleep that categorize this disease as part of the electrical status epilepticus during slow-wave sleep/epileptic encephalopathy with continuous spike-waves in sleep/epilepsy-aphasia spectrum. Since its initial recognition and characterization in 2011, there have been 13 affected males published in the medical literature representing 7 families. We have recruited an additional 14 affected males, each from a different family. The phenotypic spectrum is slightly broader than that previously reported, but there is still at least a moderate degree of developmental delay primarily affecting speech, and the seizures are poorly controlled. Continuing the trend of published cases, the presumed pathogenic variants are almost always loss-of-function (small out-of-frame deletions, splice site variants, nonsense mutations, whole gene deletions).

Conclusions CNKSR2-related disease is a severe epileptic encephalopathic syndrome with characteristic EEG patterns, poorly controlled seizures, and a developmental delay profile significantly affecting speech. The responsible molecular mechanisms continue to be almost exclusively loss-of-function variants.

265 DOCUMENTATION OF IFT172 AS THE NINETEENTH GENE CAUSATIVE OF ORAL-Facial-DIGITAL SYNDROME UNDERSCORES THE IMPORTANCE OF CIliARY TRAFFICKING IN ITS PATHOGENESIS

Yamada, Kosaki. Keio University School of Medicine, Tokyo, Japan.

Purpose of study Disruption of genes involved in the RAS/MAPK pathway is associated with well described conditions, including Noonan syndrome. However, phenotypes associated with a duplication of RAF1 are incompletely described and a unifying phenotype has not emerged. We describe two cases with duplications of RAF1. The underpinning mechanisms of this phenotype will be further explored in functional studies involving a zebrafish model.

Methods used Review of medical records and literature; over-expression ofraf1 in zebrafish model.
Summary of results The first individual is a 4yo female who presented in infancy with hypertonia, developmental delay, small PFO, pseudoarthrosis, hypertelorism, mild ptosis, mild pectus excavatum, and plagiocephaly. Over time, gross motor delay improved but speech delay persisted. Chromosomal microarray showed a 921 kb duplication resulting in a trisomy of 3p25.2, which included RAF1 in addition to SYN2, TIMP4, PPARG, TSEN2, MKRN2OS, MKRN2, TMEM40, CAND2, RPL32, SNOQATA and IQSEC1. The second individual is a 13mo male with neonatal feeding issues and respiratory difficulties requiring mechanical ventilation. CT-angiogram with 3D reconstruction identified an aberrant subclavian artery. He underwent surgery and g-tube placement. Clinical features included short stature (<3rd centile), short neck, low hairline, and minor facial anomalies. Chromosomal microarray identified a small duplication (180kb) including only the RAF1 and TMEM40 genes. Overexpression of wild type RAF1 in a zebrafish model is underway and results will be presented.

Conclusions We present two patients with microduplications that include RAF1, but neither exhibited a typical Noonan syndrome phenotype. The findings of these two patients suggest that increased gene dosage of RAF1 results in phenotypes that do not completely exhibit all of the classic Noonan syndrome features. To date, no functional studies have been performed to determine the impact of a duplication of RAF1 and whether there is a reproducible constellation of clinical findings. Results of the effects of RAF1 overexpression in zebrafish from pending studies will help provide additional information.

Purpose of study As children become ambulatory, the incomplete development of motor and balance skills may predispose them to injury. These injuries are less likely to present in pre-ambulatory children and may indicate a genetic condition, metabolic disorder, or non-accidental trauma. The purpose of this study was to identify fracture etiology and determine the incidence of metabolic and genetic disorders that predispose to fractures in these patients.

Methods used A retrospective chart review was completed of patients presenting to Rady Children’s Hospital San Diego and its affiliated providers between July 2011 and December 2012. Encounters for fractures in EPIC for all children ages 18 months and younger were analyzed by a team of reviewers and revised with specialty faculty if required. Etiologies were classified as Accidental, Inflicted, Genetic/Metabolic Disorder, Other Predisposing Factor, and Undetermined. Exclusion criteria included dental fractures, imaging without evidence of fracture, or imaging not associated with an encounter. Statistical analysis was completed using Stata 16 (College Station, TX). Fisher’s Exact Test was used for comparisons of demographic and clinical factors among etiologic groups. Cases of Undetermined etiology were excluded from group-wise comparisons.

Summary of results 696 encounters were analyzed with a mean presenting age of 10.6 months (range: 3 days to 18 months). Etiologies of fractures by research team review were as follows: 516 cases (85.1%) of Accidental injury; 30 (5.0%) of Inflicted injury; 5 (0.8%) due to a Predisposing Factor; 4 (0.7%) due to a Genetic Disorder known to predispose to fracturing; and 51 (8.4%) of Undetermined cause. The Genetic Disorders included hypophosphatasia, osteopetrosis, osteogenesis imperfecta, and hereditary sensory and autonomic neuropathy. Interestingly, in 86 cases (14.2%), there was discrepancy between the initial clinical impression and the research team review of fracture etiology.

Conclusions Though the number was small, there were children with genetic disorders who presented with fractures in early age. The discrepancy between initial and review team etiologies requires further analysis into factors that may have affected these conclusions.
Clinical Genomics and Rapid Targeted Mouse Modeling: A Novel Program to Elucidate Variants and Genes of Uncertain Significance and Advance Translational Research

S Shankar*, K Grimsrud, K Lloyd, KA Rauen, UC Davis, Sacramento, CA; UC Davis, Davis, CA

Purpose of study Whole Genome Sequencing (WGS) has revolutionized gene discovery for many inherited disorders, yet majority remains elusive. We report on a novel program using whole genome sequencing and rapid targeted mouse modeling to perform in vivo functional analysis and accelerate precision medicine.

Methods used Comprehensive clinical workup including WGS in 70 individuals with Diagnostic Odysseys. Development of a novel interdisciplinary approach to in vivo modeling and functional analysis of: 1) variants of unknown significance and 2) pathogenic variants in known disease-causing genes, to advance translational research, understand disease mechanisms and develop therapeutic interventions.

Summary of results 45 of 70 WGS test results were reported: 13 positive, 23 VUS and 7 negative. Of the 11 variants chosen for mouse models, 10 have been successfully recapitulated in mice, 4 are being phenotyped, and 1 (CHAMP1) completed. The CHAMP1 variant mouse model exhibits a number of structural and behavioral abnormalities similar to the patient’s presentation, including dysmorphic craniofacial features, delayed growth, increased startle response, anxiety, and social/object novelty.

Conclusions WGS provided a definitive genetic diagnosis in 28–33% of patients; >50% of patients were reported as VUS, warranting expedited mouse modeling. Findings from in vivo functional analysis as for CHAMP1 will achieve a definitive diagnosis, contribute to clinical decision-making, and potentially accelerate implementation of an effective management plan. The UC Davis Precision Genomic Program and Mouse Biology Program have together established a process for clinicians and researchers to nominate variants for expedited production of mice, recapitulating genetic variants and, catalyzed targeted phenotyping to accelerate disease diagnosis. These small animal avatars enable rapid determination of variant pathogenicity and inform clinical case management serving to advance precision medicine.

A New Male Case of Kagami-Ogata Syndrome Caused by Paternal Unipaternal Disomy 14 Resulting from Robertsonian Translocation

H Wang*, X Wang, H Peng, B Shah, H Gu, L Zhang. Loma Linda University School of Medicine, Loma Linda, CA; The First Hospital of China Medical University, Shenyang, China; The University of Oklahoma Health Science Center, Oklahoma City, OK

Purpose of study Umbilical cord Direct Coombs (Umbilical cord Direct Coombs) is commonly used to screen for risk of hemolytic disease of the newborn (HDN) and development of hyperbilirubinemia, particularly when there is concern for ABO or Rh incompatibility between mother and neonate, or if mother is known to be antibody positive. Since 2001, our institutional policy has been to complete DCT in these scenarios and, if positive, screen the neonates’ total bilirubin (TB), complete blood count, and reticulocyte count at 12 hours of life (HOL). In 2004, our institution also started universal serum TB testing in all neonates at 18–24 HOL. The aim of this study was to analyze the utility of early testing in at risk neonates in the era of universal TB screening.

Methods used We queried the electronic health record (Epic Systems, Madison Wisconsin) for all late-preterm and term cases of HDN, only 6 have upd(14)pat from a Robertsonian translocation (ROB) and all are female. We report a newly identified male case.

Case description This naturally conceived male infant was born at 36 weeks by cesarean section due to congenital anomalies and polyhydramnios. Mother had a history of miscarriage. Family history was negative without consanguinity. BW and HC were >97th percentile with length <10th percentile. He had narrow forehead, frontal bossing, short palpebral fissures, depressed nasal bridge, anteverted nares, elongated philtrum, low-set posteriorly rotated ears, and micrognathia. Omphalocole, short neck, bell-shaped chest, elongated fifth finger with flexion contractures, deep sacral dimple, bilateral undescended testes, right hydrocele, PDA, left-sided mild hydronephrosis, 11 pairs of ribs and bell-shaped thoracic cavity with coat hanger looking ribs were identified. Chromosome analysis revealed male karyotype with a balanced ROB:45,XY,der (13;14)(q10;q10). Parental karyotyping indicated that this ROB was inherited from his father. Concern for KOS caused by upd(14)pat was raised. The subsequent microsatellite analysis demonstrated that the patient inherited two homologous chromosome 14 from his father, confirming a paternal uniparental heterodisomy of 14q. SNP array revealed two ROH and no CNV. The results are consistent with the diagnosis of KOS caused by upd(14)pat.

Discussion/Conclusion The potential mechanisms leading to UPD are multiple. Our case is considered to be trisomy rescue as the consequence of ROB. In the clinical setting, for neonates with characteristics features and ROB, further parental study and molecular analysis for UPD should be considered.
newborns discharged from the newborn nursery at University of California, San Diego Health between May 14, 2011 and May 15, 2018. Data was analyzed for presence and result of DCT, TB levels and if phototherapy was started during admission.

**Summary of results** We identified 14,764 newborns, of whom 1,051 (7.1%) received phototherapy. A total of 6,622 (44.9%) subjects had DCT and 781 had a positive result (25.0%). Of subjects with DCT, 592 (8.9%) received phototherapy (OR: 1.64, 95% CI: 1.45–1.86). Of subjects with positive DCT, 195 (25.0%) received phototherapy (OR: 4.53, 95% CI: 3.73–5.49; positive predictive value: 24.97%, 95% CI: 22.47–27.65%). In subjects who eventually required phototherapy during admission, the average TB level at 24 HOL was 7.4 mg/dL (SD 1.5) compared to 5.2 (SD 1.2) in subjects who did not receive phototherapy.

**Conclusions** In patients who qualified for DCT, there was a small, increased need for phototherapy overall, and as expected, a positive DCT resulted in an even greater likelihood for phototherapy treatment. However, the average TB level for both phototherapy and non-phototherapy groups at 24 hours of life did not meet treatment thresholds based on the American Academy of Pediatrics guidelines. Therefore, 24 HOL screening was acceptable to help risk stratify patients, and escalation of screening of TB at 12 HOL may not be essential in the setting of universal TB screening.

**Conclusions** Neonates with Down syndrome have increased risk of early hyperbilirubinemia and need for phototherapy when compared to controls. Clinicians caring for neonates with Down syndrome are advised by the AAP to obtain an early screening CBC. We submit that simultaneously obtaining a TSB is also advisable.

### Abstract 273

**IMPACT OF A PARENT VIDEO VIEWING PROGRAM IN THE NEONATAL INTENSIVE CARE UNIT**


1. University of Utah, Salt Lake City, UT; 2. Intermountain Healthcare, Murray, UT; 3. University of Utah and Salt Lake City VA Hospital, Salt Lake City, UT; 4. Stanford University, Stanford, CA

#### Purpose of study

To compare total serum bilirubin levels, phototherapy usage, and hospital readmission for jaundice treatment, among neonates with Down syndrome vs. controls.

#### Methods used

We collected first total serum bilirubin (TSB) values, phototherapy administration, and readmission for jaundice on neonates ≥35 weeks gestation born at Intermountain Healthcare hospitals from 2004 through 2018. We created reference intervals for TSB during the first 48 hours after birth, from control neonates who did not have Down syndrome, a positive DAT, an eventual diagnosis of G6PD deficiency, or hereditary spherocytosis. We then determined the proportion of neonates with Down syndrome whose TSB exceeded the 95th percentile control interval, the percentage receiving phototherapy in each group, and the rate of readmission to the hospital for jaundice treatment in each group.

#### Summary of results

We compared 357 neonates with Down syndrome with 377,368 controls. Those with Down syndrome had 4.7 times the risk (95% CI: 3.9–5.7; p < 0.0001) of an initial TSB exceeding the 95th percentile control interval, 8.9 times (95% CI: 8.1–9.8; p < 0.0001) the phototherapy usage, and 3.6 times (95% CI: 1.6–8.2; p = 0.0009) the readmission rate for jaundice treatment.

**Abstract 273 Table 1** Survey responses for variables of interest

<table>
<thead>
<tr>
<th>Question</th>
<th>Video S1</th>
<th>Video S2</th>
<th>No video S1</th>
<th>No video S2</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>I plan to breastfeed</td>
<td>7 (7–7)</td>
<td>7 (7–7)</td>
<td>7 (6–7)</td>
<td>6 (3–7)</td>
<td>0.01</td>
</tr>
<tr>
<td>I feel involved in my baby</td>
<td>6 (5–7)</td>
<td>7 (7–7)</td>
<td>6 (6–7)</td>
<td>7 (6–7)</td>
<td>0.08</td>
</tr>
</tbody>
</table>

Median (IQR): Video N=30 and No video N=70.
TEACHING IN THE NICU: CAN WE DO BETTER?
S Langston*, A Relan, VP Walker, J Enciso, J Lloyd, M Nguyen. UCLA, Los Angeles, CA
10.1136/jim-2019-WMRC.274

Purpose of study A needs assessment revealed pediatrics interns received variable teaching on core neonatology topics. We proposed a flipped classroom (FC) approach to promote active learning through engagement in case-based discussions. FC may improve knowledge acquisition and increase pediatric trainee competency with management of critically ill neonates.

Methods used The intervention involves pediatric interns rotating through one Level III NICU. Interns complete pre-surveys of self-reported competency and knowledge in FC and non-FC topics prior to the rotation as they serve as their own matched control. Competency is scored on a scale of one to five, with five being the most competent. During their two-week rotation, interns review six brief, online digital presentations addressing core NICU topics. All six topics are discussed in the form of case-based learning in the NICU. Interns are assessed on other critical NICU topics not included in the FC curriculum to compare the impact of FC knowledge acquisition and competency with traditional teaching. Instruction of these additional subjects is at the discretion of the attending neonatologist. At the end of the rotation, interns complete the same surveys of competency and knowledge to assess change.

A general evaluation to assess satisfaction with the FC curriculum is completed.

Summary of results Scores are reported as means ± standard deviation. 13/14 interns agreed the FC approach improved decision-making, but 11/14 reported no preference for the FC curriculum over didactics. Matched-pairs T-Test showed average change in self-reported competency was greater for FC (10.8±2.93) than non-FC topics (5.4±4.23) (p<0.01). Knowledge scores showed an increase from start (58.91±16.56) to end (56.27±6.82) of the rotation for all topics (p<0.01). At the end of the rotation, the number of FC questions missed was lower than non-FC topics missed (p<0.01). Non-FC topics were inconsistently taught in the NICU. Overall, residents reported positive satisfaction levels with FC.

Conclusions Our study demonstrates potential efficacy of a FC approach to promote knowledge and competency among physicians-in-training. Pediatric interns reported positive satisfaction levels. Use of FC throughout other training programs may enhance graduate medical education.

HIDDEN COSTS OF TIME: A COST COMPARISON OF CREATING VERSUS UPDATING EDUCATIONAL MATERIALS IN NEONATOLOGY FELLOWSHIP
*Arias-Shah*, 1NM Carabajal, 1H Karpen, 1C Gisondo, 1M Gray, 1H French, 1S Izatt, 1L Johnston, 1P Chess, 1R Duda. Baylor College of Medicine, Houston, TX; 2Emory School of Medicine, Atlanta, GA; 3University of Michigan, Ann Arbor, MI; 4University of Washington School of Medicine, Seattle, WA; 5University of Pennsylvania, Philadelphia, PA; 6Duke, Durham, NC; 7Yale School of Medicine, New Haven, CT; 8University of Rochester, Rochester, NY
10.1136/jim-2019-WMRC.275

Purpose of study The flipped classroom (FC), compared to traditional didactics (TD), is an increasingly popular educational method. The Organization of Neonatal-Perinatal Medicine Training Program Directors (ONTPD) National Neonatology Curriculum (NNC) consists of peer-reviewed educational materials for FC learning of neonatal physiology and is free to fellows and educators. Potential cost savings of shared national resources for education materials is largely unstudied. We aim to: (1) define the costs of creating versus updating existing TD materials (data presented below) and (2) perform a cost comparison of TD and FC methodologies (data collection in process).

Methods used As part of a randomized controlled trial comparing the effectiveness of TD versus FC methods, an IRB-approved survey was sent to faculty querying the time required to create versus update pre-existing TD materials and costs were calculated accordingly. A follow-up survey querying the time required to develop FC materials will be evaluated similarly and compared with TD data.

Summary of results Ninety-six percent (298/309) of educators responded. The time spent creating new TD materials was significantly different from updating pre-existing materials (6 hrs vs 1 hr, P >0.0001). There was a four-fold difference in the cost to develop versus update TD materials ($692.28 vs $115.38), with a higher cost per learner in smaller fellowships. Similar calculations will be conducted for developing FC material.

Conclusions There is a greater cost associated with creating TD materials compared to updating existing materials, and the observed difference is highest in smaller-sized fellowships. A detailed analysis of how time is spent on developing FC materials for the NNC can provide insight as to how nationally shared peer-reviewed educational materials can potentially decrease the overall cost of education amongst fellowship programs.

REDUCING LENGTH OF STAY IN NEONATAL ABSTINENCE SYNDROME THROUGH QUALITY IMPROVEMENT
R Carter*, K Weiss, R Manrique, C Le, S Fleming, G Romanowski, D Golembeski, R Song, L Moyer. Rady Children’s Hospital, San Diego, CA
10.1136/jim-2019-WMRC.276

Purpose of study Neonatal abstinence syndrome (NAS) is characterized by abrupt drug withdrawal at birth in infants exposed to chronic in-utero opioids, and its frequency over the past decade has tripled due to rising opioid use and misuse. The use of standardized protocols for treatment of NAS can optimize outcomes, reduce opioid exposure, and reduce length of stay (LOS). Our current setting is a hospital system with seven Level II-IV NICUs caring for patients with NAS. There is significant variability and lack of standardization in treatment for our NAS population with an elevated length of stay. Our specific aim is to reduce LOS in infants with NAS from 22 days to 18 days by June 2020.

Methods used A multidisciplinary taskforce was established and monthly meetings with nursing, physicians, pharmacy, and ancillary staff were held. Baseline data on LOS for infants with NAS requiring pharmacologic treatment were collected from all sites. LOS for infants requiring pharmacologic treatment are tracked as a primary outcome measure.
Abstracts

277 ROLE OF ALPHA-1 RECEPTORS IN THE CEREBRAL BLOOD FLOW RESPONSE TO ACUTE HYPOXEMIA IN FETAL SHEEP

1L. Bierwirth*, 1T. Liu, 1B. Mendez, 2G. Bragg, 3G. Power, 1, 2A. Blood. 1Loma Linda University School of Medicine, Loma Linda, CA; 3Longo Center for Per. Bio., Loma Linda, CA

Purpose of study Acute hypoxemia is a common fetal stressor during gestation and labor. In response, the fetus increases cerebral blood flow (CBF) and decreases flow to the periphery. During systemic hypoxemia, peripheral vasoconstriction is mediated by catecholamines acting on alpha-1 receptors (a1R). a1R are also present in cerebral arteries where they mediate vasodistraction. We hypothesized that a1R in the fetal brain counteract increases in CBF during acute hypoxemia.

Methods used Near term fetal sheep (0.9 gestation) were chronically instrumented with central arterial and venous catheters and a composite laser doppler flow/tissue PO2 (tPO2) probe in the parietal cortex. After 4 days of recovery, by lowering the ewe’s FiO2, fetuses underwent 40-min of isocapnic hypoxemia (FiO2 reduced from 0.21 to 0.10). CBF and tPO2 were continuously recorded and arterial blood gas samples collected.

Conclusions Blocking of a1R during acute hypoxemia did not result in increased CBF or decreased CVR, suggesting that a1R do not counteract increased CBF during hypoxemia. CVR is elevated following hypoxemia and this is not mediated by a1R.

278 IUGR, DEVELOPMENT, AND DHA SUPPLEMENTATION ALTER RAT HEPATIC PEMT EXPRESSION

S. Kasera*, B. Zhao, H. Wang, L. Joss-Moore. University of Utah, Salt Lake City, UT

Purpose of study Intrauterine growth restriction (IUGR) reduces neonatal circulating docosahexaenoic acid (DHA), an ω-3 fatty acid essential for organogenesis and favorable neonatal outcomes. Mechanisms by which IUGR decreases circulating DHA are not completely understood. However, one potential mechanism involves hepatic metabolism of DHA and other long-chain fatty acids by the phosphatidylethanolamine methyltransferase (PEMT) phospholipid methylation pathway. Impaired PEMT activity reduces circulating DHA by increasing hepatic DHA accumulation in the form of phosphatidylethanolamine (PE) containing lipid droplets. We previously demonstrated that IUGR increases hepatic lipid droplet accumulation in weaning male, but not female, rat pups. However, the temporal changes in hepatic PEMT expression and the effects of IUGR and DHA supplementation on hepatic PEMT are unknown.

We hypothesized that IUGR increases hepatic lipid droplet accumulation in weaning male, but not female, rat pups. However, the temporal changes in hepatic PEMT expression and the effects of IUGR and DHA supplementation on hepatic PEMT are unknown.

Methods used We measured mRNA in control rat liver at birth and day 4, 7, 10, 12, 14, and 21 after birth, using real-time RT PCR. IUGR rats received a regular diet or a 0.1% DHA diet throughout pregnancy and lactation. At postnatal day 21 (D21), hepatic PEMT mRNA and protein were measured using real-time RT PCR and western blot.

Summary of results Both male and female IUGR rat pups weighed less than sex-matched control pups at birth and at D21. Hepatic PEMT mRNA levels were detectable from birth (MAP), heart rate (HR), CBF, cerebral vascular resistance (CVR) and tPO2 were continuously recorded and arterial blood gas samples collected.

Conclusions Lowering the FiO2 in the ewe reduced uterine artery blood flow (CBF) and decreased PO2 in the fetal lamb. Hypoxemia during gestation and labor is associated with increases in cerebral blood flow (CBF) and decreases in peripheral blood flow. In response, the fetus increases CBF and decreases flow to the periphery. During systemic hypoxemia, peripheral vasoconstriction is mediated by catecholamines acting on alpha-1 receptors (a1R). a1R are also present in cerebral arteries where they mediate vasodistraction. We hypothesized that a1R in the fetal brain counteract increases in CBF during acute hypoxemia. CBF and tPO2 were continuously recorded and arterial blood gas samples collected.

Conclusions Blocking of a1R during acute hypoxemia did not result in increased CBF or decreased CVR, suggesting that a1R do not counteract increased CBF during hypoxemia. CVR is elevated following hypoxemia and this is not mediated by a1R.

Conclusions Blocking a1R during acute hypoxemia did not result in increased CBF or decreased CVR, suggesting that a1R do not counteract increased CBF during hypoxemia. CVR is elevated following hypoxemia and this is not mediated by a1R.
Bronchopulmonary dysplasia (BPD) continues to be a significant contributor to morbidity and mortality in premature infants. Despite improvements in neonatal care, the ability to reliably predict BPD development remains limited, especially early in clinical course.

Machine learning can be used to build predictive models that outperform traditional predictive approaches. Here we use Gradient Boosting (GB), a powerful ensemble machine learning algorithm, to predict BPD development on postnatal day one of life.

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Machine learning can be used to build predictive models that outperform traditional predictive approaches. Here we use Gradient Boosting (GB), a powerful ensemble machine learning algorithm, to predict BPD development on postnatal day one of life.

Methods used This is a single center [Harbor-UCLA Medical Center Neonatal Intensive Care Unit (NICU)] study, 60 variables available within the first 24 hours of life in the California Perinatal Quality Care Collaborative database were extracted on all admissions to Harbor-UCLA NICU from 2005–2018. Variables missing data for more than 25% subjects were excluded. Infants with a major congenital anomaly (n=325) were also excluded. A total of 1832 infants were included for analysis.

A GB model was developed using the pandas, scikit-learn and XGBoost libraries in the Python programming language. Hyperparameter optimization to increase model performance was done using Exhaustive Grid Searching. Stratified K Fold (K=20) cross-validation was performed to train and test the model. Model performance was evaluated from the resultant confusion matrix.

Summary of results For the trained and cross-validated model, accuracy was 91.1%, Sensitivity was 70.3%, and specificity was 93.4%. Area Under Receiver Operating Curve (AUROC) was 0.944. The accuracy and AUROC of our model exceeded that of a logistic regression by around 4.5% and 0.082, respectively. According to the model, variables that contributed most towards the final prediction (maximal information gain) were surfactant administration, post-delivery room high frequency ventilation, and postnatal steroids.

Conclusions Our GB model yielded a high specificity for prediction of BPD in a heterogeneous population of infants. These data can be utilized for risk stratification and prognostication even on postnatal day one of life.
require a micronutrient such as selenium (Se) or zinc (Zn) to function at maximum efficacy. These trace elements are low in neonates compared to adults.

The liver is an important immune surveillance organ where antioxidant defense is critical for host response. It also plays a major role in micronutrient processing. However, the developmental regulation and expression of AOE in the liver is incompletely described. We hypothesized the neonatal liver would have decreased Se and Zn associated AOE.

Methods used C57BL/6 mice were sacrificed at P0, P7, P21 and 8–12 weeks. mRNA and protein expression for key AOE (SOD1, SOD2, SOD3, Gpx1, Gpx4, Mrsb1, TrxR1) and factors for Se processing (Sehps2/Sps2, Scly) were measured by qPCR and Western blot. Activity level pending.

Summary of results Hepatic selenoenzymes Gpx1 and Mrsb1 were developmental regulated, with mRNA low at P0 and increased by adult (p<0.05, n=5–6). Gpx1 protein increased 9-fold and Mrsb1 protein increased 6-fold from P0 to adult (p<0.0001, n=2–4). Gene expression of Zn related SOD1 and SOD2 increased postnatally, low at P0 and increased in adult (p<0.01, n=5–6). Protein expression for each increased 3-fold from P0 to adult (p<0.001, n=4). The mRNA and protein expression for Gpx4, TrxR1, TrxR2 and SOD3 remained constant postnatally.

As the greatest induction was observed in selenoenzymes, factors for Se processing were evaluated. Sehps2 and Scly mRNA increased from P0 compared to P21 and adult (p<0.05, n=4–6). Protein expression for Sps2 and Scly increased postnatally (p<0.01, n=4).

Conclusions The liver experiences a postnatal induction in essential Se and Zn associated AOE. Additionally, the hepatic machinery for Se processing is low in neonatal mice. We speculate that the neonatal liver is vulnerable to oxidative stress secondary to low AOE defense. We also speculate states that decrease neonatal micronutrient status may further impair the hepatic redox state.

**DYSRUPTION OF CALCIUM HOMEOSTATIS AMPLIFIES TLR9 MEDIATED IL-1 EXPRESSION: IMPLICATION FOR STERILE INFLAMMATION**

R de Dios*, L Nguyen, S McKenna, J Sandoval, C Wright. University of Colorado Anschutz Medical Campus, Aurora, CO

Purpose of study Sterile inflammation and IL1 expression is central to the pathogenesis of ischemia reperfusion injury. Mitochondrial stress is thought to be involved in initiating sterile inflammation. With injury, CPG rich mitochondrial DNA leaves the mitochondria and acts as a TLR9 ligand initiating pro-inflammatory signaling. Studies have used synthetic Cpg oligodeoxynucleotides to study this process; however, mitochondrial stress also disrupts calcium homeostasis. How this affects CPG-mediated TLR9 activation is unknown. Understanding the relationship between calcium and CPG-mediated TLR9 activation could identify therapeutic targets to reduce tissue injury linked to sterile inflammation.

Methods used RAW 264.7 macrophages were exposed to CpG DNA (3 μM 0–4 hours) to interrogate innate immune signaling. To assess intracellular calcium effects on CPG mediated innate immune signaling, RAW 264.7 were exposed to CpG DNA (3 μM 0–4 hours) and a) calcium ionophore A23187 (10 μM) b) calcium chelator EGTA/AM (250 μM) or c) calcineurin inhibitor FK-506 (10 μM). TLR9 mediated NFkB activation was assessed by cytosolic degradation of NFkB inhibitory proteins and NFkB subunit nuclear translocation by Western blot. IL1α and IL1β expression was assessed by qPCR.

Summary of results Exposing RAW 264.7 to CpG led to cytosolic degradation of IkB NFkB inhibitory proteins, NFkB subunit nuclear translocation, and increased IL1α and IL1β expression. Exposure to calcium ionophore significantly increased CPG-induced IL1α and IL1β expression which was associated with enhanced degradation of NFkB inhibitory proteins. In contrast, both EGTA/AM and FK-506 with CpG DNA led to decreased expression of CPG-induced IL1α and IL1β and was associated with attenuated CPG-mediated degradation of IkB NFkB inhibitory proteins.

Conclusions A combination of CpG DNA release and increased intracellular calcium accelerates TLR9 signaling which synergistically increases IL1α and IL1β expression causating an increased inflammatory response. Based on these results, the targeting of disrupted calcium homeostasis in addition to CpG DNA poses an interesting opportunity for synergistic therapies to reduce tissue injury linked to sterile inflammation.

**Surgery III**

**Concurrent session**

**8:00 AM**

**Friday, January 24, 2020**

**283 RISK FACTOR ASSESSMENT ANALYSIS FOR PATIENTS UNDERGOING BREAST CANCER RECONSTRUCTION TO ENHANCE RECOVERY**

CS Drew*, C Lee, L Tobing, S Roddick, S Gupta. Loma Linda, Loma Linda, CA

Purpose of study Rates of mastectomy followed by breast reconstruction as a treatment plan for breast cancer have risen significantly in the past few years. However, patients undergoing immediate reconstruction at the time of mastectomy tend to have a much higher rate of complications. Variables such as history of hypertension, depression, anxiety, diabetes mellitus, obesity, asthma and anemia have multiple systemic consequences and could potentially contribute to a higher risk of complications in this population. This study aims to better identify possible risk factors that may predispose a patient to complications post breast reconstruction.

Methods used This is a retrospective study that collected data from July 2014 to June 2019 of patients who underwent breast reconstruction in a single institution. Comorbidities were gathered and statistical analysis was performed to analyze commonalities shared between patients predisposing them to higher rates of complications.
Abstract 283 Table 1

<table>
<thead>
<tr>
<th>Comorbidity</th>
<th>patients who had the comorbidity and returned with complications (n=99)</th>
<th>patients who had the comorbidity and underwent breast reconstruction (n=1817)</th>
<th>odds ratio</th>
</tr>
</thead>
<tbody>
<tr>
<td>Hypertension</td>
<td>46</td>
<td>854</td>
<td>0.979</td>
</tr>
<tr>
<td>Obesity (BMI&gt;30)</td>
<td>48</td>
<td>941</td>
<td>0.876</td>
</tr>
<tr>
<td>Anemia</td>
<td>20</td>
<td>127</td>
<td>3.369</td>
</tr>
<tr>
<td>Asthma</td>
<td>19</td>
<td>113</td>
<td>3.581</td>
</tr>
<tr>
<td>Diabetes</td>
<td>15</td>
<td>491</td>
<td>0.482</td>
</tr>
</tbody>
</table>

Summary of results The results of this study are summarized in the table below. Those with complications had a much higher incidence of anemia and asthma compared to the general population of patients undergoing reconstruction. In those with complications hypertension and obesity occurred at a similar incidence, while diabetes occurred at a lower incidence to the overall population undergoing this procedure.

Conclusions Patients undergoing breast cancer reconstruction who also had anemia and asthma had relative risks of 3.36 and 3.58 to suffer complications. In patients with complications, hypertension and obesity were observed in the same proportion as those without complications. However, diabetes mellitus was observed less often than expected in patients with surgical morbidities. This data provides additional guidance in the development of enhanced recovery after surgery protocols for the population of breast cancer reconstruction patients.

Abstract 285

ETIOLOGY OF CHRONIC EXERTIONAL COMPARTMENT SYNDROME AND A NON-SURGICAL TREATMENT


10.1136/jim-2019-WMRC.284

Purpose of study Chronic exertional compartment syndrome (CECS) occurs when increased intramuscular pressure produces severe, sometimes debilitating pain and neurologic complications. Due to incomplete understanding of the pathophysiology, both diagnostic criteria and treatments options for CECS are lacking in reliability and efficacy.

The purpose of this study is two-fold. First, to investigate the theory that CECS results from functional venous outflow obstruction due to muscular compression. Second, to investigate the treatment of CECS through an image guided botulinum toxin injection targeted at the site of functional muscular compression.

Methods used Retrospective case study of 284 patients (age 15–77, mean: 27) with exertional lower leg pain. Patients were assessed for CECS and areas of functional venous outflow obstruction using lower extremity MRI at rest and after exertion and CT Angiography at rest and with plantar flexions against resistance using the MVP Flex device.

Using both ultrasound and CT imaging guidance, a targeted injection of botulinum toxin was administered into the muscles at the site of venous compression. Patients were followed up at fixed time points over several years to track their treatment response.

Summary of results 260 of the 284 patients demonstrated both CECS and functional venous outflow obstruction on pre-treatment stress imaging. Active plantar flexion caused venous obstruction, in the absence of active CECS symptoms.

Following botulinum toxin treatment, venous compression diminished with most patients demonstrating symptom resolution and normal imaging studies. Return of symptoms occurred at an average of 6.9±2.4 months (range 2–24 months). Repeat treatments were performed in 167 patients.

Conclusions 91.5% of patients displayed both CECS and functional venous obstruction. The results of this study suggest CECS results from functional venous outflow obstruction due to muscular compression.

Targeted botulinum toxin injections at the site of venous outflow obstruction resulted in significant symptom improvement, with a short recovery period. This treatment provides a viable non-surgical alternative to treat CECS by improving venous outflow.

285 METAPHYSEAL CLAM SHELL OSTEOTOMY FOR LONG BONE DEFORMITY

1. DA Brainin, 2. M Rice*, 1. MA Lee. 1. UC Davis Health, Sacramento, CA; 2. UC Davis, Sacramento, CA

10.1136/jim-2019-WMRC.285

Purpose of study Metaphyseal malunions and nonunions pose a challenge especially in the setting of complex deformity. Common options have significant limitations, especially in poor quality bone and when deformity centers are near the joint. The clamshell osteotomy has been described as treatment for complex diaphyseal malunions and provides efficient acute multiplanar correction and allows for early weight bearing. The purpose of this study is to report union and complication rates of metaphyseal and metadiaphyseal clamshell osteotomies.

Methods used After obtaining IRB approval, patients were identified using CPT codes from 01/01/2010 to 09/30/2018. Inclusion criteria were age >18 and a metaphyseal or metadiaphyseal clamshell osteotomy of the tibia or femur, indicated by malunion or nonunion. Exclusion criteria were <6 mos of follow up. Chart and radiograph review was used to determine time to union and complications.

Summary of results 15 patients met inclusion criteria. 12 of 15 patients achieved union by 16 mos (union rate of 80%). 10 of 15 (66.7%) patients healed after the primary procedure, at
an average of 8.75 mos. of 15 (26.7%) patients required a secondary procedure (exchange intramedullary nailing). 2 of these 4 patients went on to union, while the other 2 did not have union at final follow up. 1 of 15 (6.7%) patients was lost to follow up at 8 mos without union. 2 patients had a post-op PE, 1 had a draining wound that required a sinus tract excision, and 1 developed a non-displaced fracture from the RIA.

Conclusions The clampshell osteotomy is one potential tool for treating complex multiplanar deformities in the meta-diaphyseal and metaphyseal portion of lower extremity long bones. A high union rate can be seen using modern nail designs and atraumatic technique; however, union times can be long and some patients will require secondary procedures.

286 DETERMINANTS OF CAREGIVER SATISFACTION IN PEDIATRIC ORTHOPEDICS

1M Singleton*, 2R Garfinkel, 3H Temkit, 4MV Belthur. 1University of Arizona College of Medicine – Phoenix, Phoenix, AZ; 2Henry Ford, Detroit, MI; 3Phoenix Children’s Hospital, Phoenix, AZ

Purpose of study With healthcare transitioning from volume-based to value-based, patient satisfaction is becoming increasingly tied to physician reimbursement as well as publicly reported. Aside from its use as a quality metric, it is also a key component of patient-centered care. This study investigates determinants of pediatric orthopedic patients’ parent or guardian (caregiver) satisfaction with the provider as measured by the Clinician and Group Consumer Assessment of Healthcare Providers and Systems (CG-CAHPS).

Methods used This was a prospective cross-sectional study of 200 English-speaking caregivers of pediatric patients that checked into the Phoenix Children’s Hospital orthopedic clinic from March 1, 2017 to November 1, 2018. All patients saw the same attending physician. Questionnaires given in clinic included the Newest Vital Sign (NVS) and the Literacy in Musculoskeletal Problems (LiMP) survey to measure general and musculoskeletal health literacy, respectively, demographic information, wait time, Consultation and Relational Empathy Measure (CARE) to measure perceived physician empathy, and CG-CAHPS.

Summary of results Of the factors measured, perceived physician empathy correlated the strongest with the caregiver’s overall physician satisfaction. Pearson correlation coefficient yielded an r of 0.740 and a p-value of <0.0001. Using multi-variable modeling, physician empathy alone accounted for 53% of the variation in satisfaction scores. Other factors such as health literacy as measured by the NVS (r=0.004; p=0.964), LiMP (r=0.013; p=0.879), wait time (r=-0.003; p=0.974), and time spent with the physician (r=0.016; p=0.866) did not independently affect satisfaction.

Conclusions The main determinant of caregiver satisfaction with the provider in a pediatric orthopedic setting is perceived physician empathy, accounting for over half of the variation in satisfaction. Health literacy, wait time, and time spent with the physician do not significantly affect satisfaction. This highlights the importance of quality over quantity patient-physician interactions. This is the first study, to our knowledge, that directly correlates pediatric caregivers’ perceived physician empathy with provider satisfaction.

287 DOES ULTRASOUND-GUIDED OSTEOTOMY IMPROVE THE AESTHETIC OUTCOME OF PATIENTS UNDERGOING RHINOPLASTY SURGERY?

C Lee*, CS Drew, L Tobing, S Roddick, S Gupta. Loma Linda University, Loma Linda, CA

Purpose of study General ultrasound imaging is a widely available, easy-to-use, and cheaper alternative to other advanced imaging methods. Furthermore, it is noninvasive, painless, and safe as it does not have harmful radiation. However, ultrasound is not frequently used intraoperatively because of its inability to penetrate bone and large amounts of tissue to produce a clear image. In thin areas such as the nose where these restrictions can be eliminated though, this method could be highly beneficial intraoperatively by guiding nasal osteotomies during rhinoplasty to improve patients’ aesthetic outcome with enhanced visualization of symmetry.

Methods used This is a case series using ultrasound to assist osteotomies for patients undergoing nasal surgery. A SonoSite ultrasound is specifically used with the L2S 13–6 probe and under the musculoskeletal exam. The nose is seen superficially at a depth of 2.0 cm and examined longitudinally for optimal view. Gain may be adjusted for a more advantageous assessment. Ultrasound gel is placed on the patient’s nose as well as on the probe underneath the sterile wrap to provide a clear image. After confirming the bony landmarks, the plastic surgeon performs the closed osteotomy with visualization, and the ultrasound is continuously utilized to check correction and symmetry.

Summary of results Ultrasound-assisted osteotomy was used for 5 patients undergoing nasal surgery. No complications arose during the surgery. Cartilage was clearly demarcated. Seven plastic surgeons and plastic surgery residents were surveyed regarding the potential for this technology to enhance their rhinoplasty outcomes. Comments were collected from the users and observers to guide further implementation.

Conclusions Intraoperative ultrasound imaging for patients undergoing nasal surgery has so far been successful in ensuring symmetry compared to the naked eye alone. Furthermore, it has enabled surgeons to visually confirm precision and accuracy beyond the skin surface and has accordingly provided a stronger outlook on the aesthetic result for the patients. Thus, with sufficient learning and skill, intraoperative ultrasound can be a highly advantageous tool in aesthetic and reconstructive plastic surgery.

288 SELF-REPORTED HEALTH MEASURES IN BURN SURVIVORS UNDERGOING BURN SURGERY FOLLOWING ACUTE HOSPITALIZATION

1DE Luna*, 2C Schechter, 3G Carragher, 4B Stewart, 5J Schneider, 6C Ryan, 7R Holavanahalli, 8D Antmann, 9N Gibran, University of Washington, Seattle, WA; 2Stanford, Stanford, CA; 3Harvard, Boston, MA; 4UT Southwestern, Dallas, TX

Purpose of study Perception of mental and physical function in adult burn survivors is decreased following injury. Individuals who undergo post hospitalization reconstructive surgery may have lower Health Related Quality of Life (HRQoL) that is influenced by the severity of their burns. We sought to characterize self-reported health perceptions in burn survivors
undergoing surgery post-hospitalization within 24 months of injury.

Methods used Patient Reported Outcomes Measurement Information System (PROMIS)-29 Profile v2.0 data were collected from participants from a multi-center longitudinal study at 6, 12, and 24 months post-injury. Surgical operations performed within the 24-month follow-up period were classified as: wound closure, contracture release or scar revision. PROMIS-29 domain scores were compared for those undergoing surgery (surgical participants) versus those who did not (non-surgical participants) using linear regression, adjusting for age, burn size, range of motion limitation, number of operations during acute treatment, and insurance/payer.

Summary of results 727 participants provided complete PROMIS-29 data, of which 227 (31.2%) underwent >1 operative procedure within 24 months post-injury (Table). PROMIS-29 anxiety and depression scores were worse at 6 months post-injury for participants who underwent contracture release and scar revision compared to non-surgical peers (p<0.05). At 12 months after injury, individuals who underwent wound closure at any time post initial hospitalization reported worse anxiety, depression, and pain interference (p<0.05). Participants in the contracture release and scar revision categories reported lower PROMIS-29 scores in all domains 24 months after injury (p<0.05), even after adjusting for confounders.

Conclusions Burn-related reconstructive surgery has the potential to improve the lives of patients including promoting functional recovery and improving cosmetic appearance. Our data suggest that participants who undergo reconstruction surgery after hospital discharge, report worse self-reported indicators of health compared to participants who do not undergo surgery after adjusting for known confounders.
Behavior and development I

Concurrent session

10:15 AM

Friday, January 24, 2020

SPEECH AND LANGUAGE DEVELOPMENT, AUTISM SPECTRUM DISORDERS AND SEIZURES: WOULD WHOLE GENOME SEQUENCING PROVIDE A CONNECTING LINK?

SL Cole*, 1M Hegde, 1S Shankar. 1UC Davis, Sacramento, CA; 2PerkinElmer, Atlanta, GA

Purpose of study Many studies have shown a correlation between autism spectrum disorders (ASD), neurodevelopmental disorders and epilepsy/seizure disorders. We evaluated whole genome sequencing (WGS) results in a subset of patients with limited speech/nonverbal, with or without ASD, neurodevelopmental disorders and seizures to find common genetic variants/network.

Methods used Clinical whole genome sequencing was performed and relevant genetic variants were analyzed to identify recurring pathways or genes.

Summary of results We found 15 individuals in our cohort with this combination of phenotypic characteristics. Four had pathogenic or likely pathogenic variants in SLC16A2, PHF6, ERCC8 (relevant to phenotype), and MME (incidental finding), 10 patients had one or more variants of unknown significance (VUS) and 1 had a negative test result reported. VUSs included autosomal dominant inheritance in genes consistent with the phenotypes in our cohort. Additionally, some patients had multiple VUSs in causative genes suggesting oligogenic gene effects.

Conclusions WGS provides an opportunity to evaluate regions of the genome that are not picked up by traditional genetic testing. Overall, we identified many changes that have been identified as VUSs and have compensated for the variation in the parent phenotype. However, given the novelty of WGS, a direct correlation requires additional patients with WGS testing and phenotypic correlation to determine significance.

DEVELOPMENTAL DELAY AND PSYCHIATRIC ISSUES IN AN UNDER-STIMULATED DEAF CHILD

NM Shah*, Loma Linda University, Loma Linda, CA

Introduction In the United States, about 10,000 infants are born with sensorineural hearing loss, 95% of which have hearing parents. These children experience barriers to communication with their families. The lack of proper communication and stimulation in the first years of life can be detrimental to mental development. This leads to a lack of abstract thinking and problem-solving skills, ability to form...
peer relationships, and healthy self-esteem. 9% of deaf children have learning disabilities and 5% have severe developmental delays. Emotional and behavioral problems are twice as prevalent.

**Case description** A 15-year-old female born deaf and mute was brought in to the Behavioral Medicine Center (BMC) after recovering from an overdose on ibuprofen, famotidine, and ranitidine. The parents did not understand the patient’s actions and could not communicate effectively with her. They knew very little sign language and spoke mainly Spanish. The patient attends a school for deaf children and can communicate in American Sign Language (ASL) while at school. However, her interaction with family is severely limited due to both a Spanish/English communication barrier and the lack of knowledge of sign language.

Her social skills were minimal both with staff and other patients on the unit. She consistently presented as indifferent and guarded. She responded to most questions with a shrug or nod whether or not the question warranted it.

The patient showed symptoms of depression including suicidality, hypersomnia, anhedonia, low energy, decreased appetite, poor mood, and psychomotor retardation. Social workers were able to connect the family and patient to helpful mental health resources in the community. The patient was placed on Fluoxetine (40 mg) for long-term management of depression.

**Discussion** The importance of awareness campaigns for developmental health of deaf children cannot be ignored. In this case of the 15-year-old female, we see what the unfortunate effects are of lack of proper care for deaf children. Educating parents about these long-term effects may motivate them to be more proactive in caring for their deaf children. Furthermore, the support community for deaf people should be connected with the mental health community. These individuals have unique needs and life experiences that deserve careful attention in all healthcare settings.
knowledge of gender bias and sexual harassment (p=0.001) as well as their ability to confront these issues in the learning environment when they personally experience GB/SH themselves (p=0.001) or when they witness GB/SH (p=0.0001). Students also reported feelings of validation, awareness and empowerment as well as appreciation for the process of information sharing, faculty facilitation, and practical techniques gained from the workshop.

Conclusions Our study suggests that students exposed to a 120-minute workshop are better equipped to recognize and respond to GB/SH. This workshop is the first step in empowering our learners.

297 THE ASSOCIATION BETWEEN SOCIAL MEDIA USE AND THE DEVELOPMENT OF DISORDERED EATING: A CLOSER LOOK AT THE LITERATURE

Maya Hidalgo*, 2V Simhambhatla, 2P Raj, 2A Patel, 2A Saha, 2,3B Afghani. 1University of San Diego, San Diego, CA; 2UC Irvine School of Medicine, Irvine, CA; 3CHOC Hospital of Orange County, Orange, CA

Purpose of study The prevalence of social media is widespread and its effect on the etiology of eating disorders is unclear. The objective of this study is to investigate the effects of social media use on eating pathology.

Methods used A literature review utilizing PubMed and Google Scholar with the keywords Social Media Use, Instagram, Facebook, Eating Disorders, and Disordered Eating was performed. Of the 130 students (75 in SP arm, 65 in HFM arm), the SP arm showed significantly greater participation when compared to the HFM arm with 98% agreeing/strongly agreeing to have ‘actively participated’ vs. 88% (p<0.05) Additionally, students in the SP arm reported greater levels of confidence in their ability to transfer the knowledge gained into a real clinical situation with 97% vs. 89% agreeing/strongly agreeing (p<0.05). They also reported greater confidence in performing physical exams practiced during the simulation on an actual patient, with 88% vs. 80% agreeing/strongly agreeing (p<0.05). Post-simulation surveys completed by the SPs with results of the modified SEGUE showed higher ICS in the SP arm, (p<0.05).

Conclusions Our study demonstrates that during simulation training, year-3 internal medicine clerkship students felt they were more active participants, had greater levels of confidence in their ability to transfer knowledge gained into a real clinical situation, and demonstrated improved ICS when interacting with SP versus HFM. We believe this study suggests that directly interacting with SP creates a more realistic simulation experience than interacting with HFM, which leads to an improved learning environment.

Summary of results Of the 130 students (75 in SP arm, 65 in HFM arm), the SP arm showed significantly greater participation when compared to the HFM arm with 98% agreeing/strongly agreeing to have ‘actively participated’ vs. 88% (p<0.05) Additionally, students in the SP arm reported greater levels of confidence in their ability to transfer the knowledge gained into a real clinical situation with 97% vs. 89% agreeing/strongly agreeing (p<0.05). They also reported greater confidence in performing physical exams practiced during the simulation on an actual patient, with 88% vs. 80% agreeing/strongly agreeing (p<0.05). Post-simulation surveys completed by the SPs with results of the modified SEGUE showed higher ICS in the SP arm, (p<0.05).

Conclusions Our study demonstrates that during simulation training, year-3 internal medicine clerkship students felt they were more active participants, had greater levels of confidence in their ability to transfer knowledge gained into a real clinical situation, and demonstrated improved ICS when interacting with SP versus HFM. We believe this study suggests that directly interacting with SP creates a more realistic simulation experience than interacting with HFM, which leads to an improved learning environment.

Abstract 297 Table 1 Association of social media use and disordered eating

<table>
<thead>
<tr>
<th>First Author &amp; Year</th>
<th># of Subjects &amp; Age (ys)</th>
<th>Frequency and duration of social media use in subjects</th>
<th>Method of evaluation/measurement</th>
<th>Eating pathology outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Smith 2013</td>
<td>N=232; Mean age=18.72</td>
<td>Not Reported</td>
<td>The effects of maladaptive FB use on body dissatisfaction and bulimic symptoms</td>
<td>Maladaptive FB usage predicted increases in bulimic symptoms (p&lt;0.05)</td>
</tr>
<tr>
<td>Mabe 2014</td>
<td>N=44; Mean age=18.39</td>
<td>Mean time reported/FB session=20.06 min mean total time/day on FB=76.28 min</td>
<td>The relationship between FB use and disordered eating</td>
<td>Participants with the greatest disordered eating had higher FB scores p&lt;0.001</td>
</tr>
<tr>
<td>Sidani 2016</td>
<td>N=1765; Age 19-32</td>
<td>Median time/day=61 minutes median visits/week=30</td>
<td>The relationship between volume of social media use and eating concerns</td>
<td>Participants with greatest volume and frequency of use had greater odds of having eating concerns Volume: AOR=2.18 p&lt;0.001</td>
</tr>
<tr>
<td>Turner 2017</td>
<td>N=680; Mean age=24.70</td>
<td>Range between 0-450 minutes/week</td>
<td>The correlation between time spent on each social media channel and Orthorexia Nervosa symptoms</td>
<td>More time spent on Instagram is associated with greater Orthorexia Nervosa symptoms Instagram: -0.10*, FB: 0.04, Twitter2.12*, Pinterest: -0.06, and Tumblr: 0.10</td>
</tr>
<tr>
<td>Saunders 2018</td>
<td>N=637; Mean age=21.3</td>
<td>Range between 0 min-3 hours/day</td>
<td>The relationship between social networking site rating, body image, and disordered eating</td>
<td>FB, Instagram and Snapchat users demonstrated a positive relationship between comparisons, body dissatisfaction, and disordered eating, p&lt;0.005</td>
</tr>
</tbody>
</table>

FB = Facebook, p.
Conducted. Only studies published after 2012 were included and studies that did not focus on eating pathology were excluded.

Summary of results Of the 15 articles, only 5 satisfied our inclusion criteria. Each of the articles summarized in the table below incorporated a self-report survey design. Of the 5 articles, each one found a significant association between social media use and disordered eating. Three of the articles found that body dissatisfaction plays a role in the relationship between social media usage and disordered eating outcomes. Limitations of some of the studies included the self-report nature and lack of control for confounding variables leading to disordered eating.

Conclusions This review suggests a correlation between social media use and disordered eating outcomes but a causal relationship cannot be determined. Prospective controlled studies that account for confounding variables are warranted in order to make causal inferences regarding social media usage and its effects on the development of eating disorders.

Community health
Concurrent session
10:15 AM
Friday, January 24, 2020

298 ADVERSE CHILDHOOD EXPERIENCES AND POVERTY: COLLABORATIVE PARTNERSHIPS TO IMPROVE SELF-EFFICACY AND RESILIENCE IN IMPOVERISHED MINORITY YOUTH

M Cabada1, T Lee1, C Casillas1, M Chavez1, K Colwell1, R Kinman1, Fresno High School, Fresno, CA; UCSF Fresno, Fresno, CA

Purpose of study Children growing up in poverty can face a variety of adverse childhood experiences (ACES) which can negatively impact their ability to succeed in life. According to the American Psychological Association, resilience ‘is the process of adapting well in the face of adversity, trauma, tragedy, threats, or significant sources of stress’. Although impoverished adolescents want to become healthy and successful adults, they may need assistance to develop their own self-efficacy and resilience. High-risk impoverished minority adolescents thus partnered with UCSF Fresno Pediatrics and the Fresno Community Health Improvement Partnership (FCHIP) to increase self-efficacy and resilience in their fellow youth.

Methods used Students explored the effect of poverty on ACES and anonymously surveyed 155 fellow high school students about their ACES exposures. They collaborated with FCHIP’s Trauma and Resilience Network to develop their own self-efficacy and resilience, and to partner with UCSF Fresno to create an annual Health Careers Job Shadow Day to educate students on how to develop their own community action research partnerships, were the closing speakers for the 2nd annual FCHIP breakfast, and developed an interactive lunchtime session at their school to educate their fellow students on how to build resilience and decrease stress.

Conclusions Although impoverished minority youth have significant exposure to ACES, they can successfully develop community partnerships to help them identify and address issues of concern within their own communities, while helping them build resilience to mitigate the effects of these ACES on their own lives.

299 HOMELESS TO HOUSED: HOW DOES STABLE HOUSING AFFECT HEALTHCARE UTILIZATION AND PERCEPTIONS IN RESIDENTS OF THE FORGET-ME-NOT MANOR, A HOUSING FIRST PROGRAM IN JUNEAU ALASKA

N Ord*1, R Urata. University of Washington, Seattle, AK

Purpose of study Housing first (HF) is a solution to chronic homelessness demonstrated to reduce public service utilization in urban populations but less is known about rural areas. In October of 2017 Juneau Alaska, a town of 32,000 opened the Forget-Me-Not Manor to house Juneau’s most vulnerable. In five months after opening, 22 of the 32 residents experienced impressive reductions in ER visits (73% reduction), recovery center sleep off visits (99% reduction), and contacts with police officers (77% reduction) compared with 6 months prior (Heidi Brocious, UAF). However, little is known about why residents use less services and the reasons residents seek healthcare. This study examines resident’s relationship to healthcare services before and after joining HF through qualitative interviews and healthcare records analysis.

Methods used Two qualitative 30-minute interviews were conducted 3 months apart exploring resident’s definition of health, barriers to improving health, and role housing first plays in their health. Additionally, resident’s healthcare records were collected from three of the most frequently visited healthcare organizations during the period of December 1st, 2015 – December 1st, 2018. The modality of each interaction (clinic, hospital, etc.), diagnosis codes, and summary category (acute illness, maintenance of chronic disease, etc.) associated with each healthcare is recorded.

Summary of results Total participants are 17. Interviews were recorded with 17 and 15 residents respectively. Participant responses to standardized interview questions are summarized and compared in pre-defined categories. 636 Medical records are collected from 17 of 17 participants and three healthcare organizations. The data is analyzed for usage patterns by location, diagnosis codes, and summary category relative to the date residents joined housing first.

Conclusions The combination of quantitative and qualitative data is used to discuss the relationship between healthcare services and the study participants before and after joining HF. Resident’s most common diagnoses are compared with the general population and HF programs in urban populations. Differences between HF in rural and urban regions are
discussed. Limitations of this study and avenues for future research are proposed.

Purpose of study Increasing the number of underrepresented individuals in healthcare fields is critical to meeting the needs of an increasingly diverse society. The purpose of this study is to evaluate the effectiveness of an interactive health exhibit in enhancing the knowledge of youth towards healthy habits and garnering their interest towards healthcare.

Methods used Through a partnership between the UC Irvine School of Medicine and the Discovery Cube in Santa Ana, CA, we have established an interactive exhibit where trained volunteers teach the young visitors about the hazards of smoking using models, introduce them to basic life support, and teach them intubation using a simulator. A feedback survey is distributed to the visitors for any qualitative feedback.

Summary of results 526 feedback surveys were collected since 2016. When visitors were asked if their interest in healthcare had increased, 378 (72%) responded and 28% left it blank or did not know. Of 378, 279 (74%) gave a positive response. When asked what they liked most about the exhibit, 103 (61%) of the 170 underrepresented (UR) and 181 (55%) of the 327 non-underrepresented (NUR) provided comments (Table). Of 284 respondents, 33% liked the hands-on nature, 28% liked the knowledge gained, and 27% liked the interaction with knowledgeable mentors. Interestingly, the underrepresented were more likely to comment that their knowledge had increased (UR=39% vs NUR=23%) and the non-underrepresented were more likely to enjoy the interactive nature of the exhibit (NUR=37% vs UR=25%).

Conclusions Our interactive exhibit was effective in garnering the interest of a diverse population of youth towards healthcare. Such interactive models can be used to enhance the knowledge and stimulate the interest of youth in school and non-school settings.

Abstract 300 Table 1 Results of survey based on themes

<table>
<thead>
<tr>
<th>Themes</th>
<th>N=181</th>
<th>UR, N(%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Interactive and Demonstrative Models</td>
<td>67 (37%)</td>
<td>26 (25%)</td>
<td>p=0.0487</td>
</tr>
<tr>
<td>Increased Scientific Knowledge</td>
<td>41 (23%)</td>
<td>40 (39%)</td>
<td>p=0.004</td>
</tr>
<tr>
<td>Engaging and Knowledgeable Mentors</td>
<td>53 (29%)</td>
<td>25 (24%)</td>
<td>p=0.41</td>
</tr>
<tr>
<td>Increased Health Awareness</td>
<td>18 (9%)</td>
<td>9 (9%)</td>
<td>p=0.83</td>
</tr>
<tr>
<td>Ideal Environment</td>
<td>5 (2%)</td>
<td>3 (3%)</td>
<td>p=1.0</td>
</tr>
</tbody>
</table>

NUR = Non-Underrepresented, UR = Underrepresented

Abstract 301 Table 1 Displays awareness percentages of subgroups

<table>
<thead>
<tr>
<th></th>
<th>Total</th>
<th>Gender</th>
<th>Language</th>
</tr>
</thead>
<tbody>
<tr>
<td></td>
<td></td>
<td>Female</td>
<td>Male</td>
</tr>
<tr>
<td>Tdap &amp; Flu Vaccine</td>
<td>32.4%</td>
<td>40.4%</td>
<td>22.1%</td>
</tr>
<tr>
<td>p-Value</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Tdap Vaccine Only</td>
<td>40.3%</td>
<td>54.2%</td>
<td>22.1%</td>
</tr>
<tr>
<td>p-Value</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
<td>&lt;0.001</td>
</tr>
<tr>
<td>Flu Vaccine Only</td>
<td>80.9%</td>
<td>75.3%</td>
<td>88.2%</td>
</tr>
<tr>
<td>p-Value</td>
<td>&lt;0.005</td>
<td>&lt;0.005</td>
<td>&lt;0.005</td>
</tr>
</tbody>
</table>

Conclusions Overall, vaccine awareness was low, driven mostly by ignorance of the Tdap vaccine, as more participants knew of the flu than Tdap vaccine. More women than men were aware of the Tdap vaccine, whereas men were slightly more aware of the flu vaccine. Nearly double the number of women were aware of both vaccines than men. Additionally, Spanish speakers were less aware of the Tdap vaccine than English speakers and awareness of both vaccines by Spanish speakers was less than a third of that of English speakers. To increase vaccine awareness and decrease infection, recommendations for pregnant women to encourage their partners and baby’s contacts to receive both vaccines should be made. Also, vaccine education initiatives should be enhanced for Spanish-only speaking populations.
AN ASSESSMENT TO REDUCE SMOKING DURING PREGNANCY IN PARK COUNTY, WYOMING

A Golden*, University of Washington School of Medicine, Seattle, WA

Purpose of study Powell, Wyoming is an agricultural community with a mean household income and education level below the national average. Through interviews with local physicians and community organizations, a lack of education and resources for mothers were found to be public health concerns. Specifically, smoking during pregnancy was identified. Data collected found that 16.1% of women in the county smoked during their pregnancy, which is above the national average.

Methods used An asset-based approach was used to determine available resources for mothers in Powell. The Serenity Pregnancy Resource Center and the Public Health Office were identified as organizations providing education and services for mothers. Using information gathered from community-based interviews, a literature review was conducted to identify possible ways to reduce smoking in pregnant women using the resources available within the town.

Summary of results Five literature review articles that proposed interventions to help with smoking cessation during pregnancy were evaluated for their potential efficacy in Powell. One of the most promising interventions proposed initiating a text-message based support group for mothers. An additional study evaluated the use of smoking cessation therapy in pregnancy. Both of these studies have realistic uses in a rural setting and are able to use resources and assets that already exist in Powell. The Serenity Pregnancy Resource Center provides education for mothers and already has a database that could be used for the text messaging program. The Public Health office provides many services to pregnant and new mothers and could potentially implement a medication cessation service.

Conclusions The assessment to help pregnant women quit smoking was aided through the use of a strength-based framework. Available resources within the community were identified and information about this public health concern were shared with local providers and community organizations. Next steps include establishing a Mother Child Health Coalition with the Public Health Office. The development of this coalition has been initiated and is pending input from providers in Park County who treat pregnant women and new mothers. Maternal smoking will be one area of focus and evidence-based interventions from this research will be employed before and during official implementation of the program.

SUPPORTING SAFE AND SUSTAINABLE EMPLOYMENT AMONG RESIDENTS IN PROTECTED HOMES FOR PEOPLE WITH SERIOUS MENTAL ILLNESS IN IQUITOS, PERU

A Larson*, University of Washington, Seattle, WA

Purpose of study As part of a national reform towards community-based mental health services, a network of protected homes for people with serious mental illness were introduced in Iquitos, Peru starting in 2012. While these homes are intended to serve a transitional role, challenges in obtaining employment and independent living for the residents have persisted since the protected homes were first established. Staff in one home proposed the introduction of an employment program in the form of a self-maintaining shop within one of the homes. However, there was initially no evaluation tool specifically for monitoring residents’ progress in occupational capacities, and tools to promote workplace health and safety as well as best practices were also missing.

Methods used A survey focused on occupational development was developed, using a variety of existing tools, for ongoing evaluation of residents participating in the employment program. An informal health and safety assessment for use by staff was developed based on tools from PAHO and the Peruvian government and piloted with staff and residents in the shop location. In addition, a team-based best practices checklist was developed for administration of occupational programs for people with serious mental illness was developed, based on existing literature.

Summary of results After feedback from staff and residents, the evaluation survey was streamlined into 5 short domains including Employment Functioning, Role and Identity, Self-Efficacy, Skills, and Quality of Life. The survey used a consistent numerical scoring system throughout, with optional prompts for further discussion between the evaluator and resident. The Health and Safety checklist was successfully employed with staff at the planned project site and used to inform the introduction of safety infrastructure in the home. The Best Practices checklist was reviewed and approved by staff on multiple levels.

Conclusions The employment program is equipped with tools to evaluate progress of individual residents, conduct ongoing internal reviews of workplace health and safety, and promote best practices through team discussion. These tools should be employed before and during official implementation of the program.

COMMUNITY-BASED STRATEGIES TO REDUCE DIABETES INCIDENCE THROUGH EDUCATIONAL GARDENING IN GRAND COULEE, WASHINGTON

E Kershaw*, University of Washington, Spokane, WA

Purpose of study Grand Coulee, WA is a rural community located in a USDA food desert. It is home to Coulee Medical Center (CMC), a critical access hospital that serves 7,000 patients from surrounding counties including Lincoln, Douglas, Grant, and Okanogan and has a 21% Native American patient population. These counties have obesity rates 2-7% higher than the state average. Community conversations revealed a concern for obesity in young people. Health care providers cited diabetes management adherence concerns, which includes lifestyle modifications such as obesity. One primary factor contributing to obesity is unhealthy food choices, with selected secondary risk factors including lack of healthy food access and limited healthy food knowledge.

Methods used Using an asset-based approach, community resources working to address secondary risk factors were evaluated. The Care and Share food bank and the local
Senior Center Meals program identified less nutritious food choices in a free food setting. For example, the selection of white bread rather than multigrain bread. Local individuals cited high food prices in general as a concern. Food education is currently addressed at CMC and the local senior center by CMC staff. The Chamber of Commerce has established a plot of land which can be utilized for a community garden. Therefore, community gardening strategies and their efficacy in improving obesity risk was the literature review focus.

**Summary of results** A successful clinic-based community gardening program includes provider encouragement, nutritional and gardening-based educational components, and utilization of volunteers. Such programs have been shown to be a strong educational addition to communities, with increases in quantity of purchased produce, regardless of income. Community gardening itself will not alleviate food insecurity, but can serve to strengthen awareness of the local food bank and senior center, which aim to meet this need.

**Conclusions** This project highlights how a community’s concerns can be addressed by utilizing its strengths. Next steps include lease of land by CMC, gardening expert-informed establishment of layout and rules, garden promotion, and securing funding for supplies via food desert-based grant funding and/or local business donations.

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**Abstract 305 Table 1**

<table>
<thead>
<tr>
<th>Variable</th>
<th>P-value</th>
<th>OR</th>
<th>95% CI</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age</td>
<td>0.0927</td>
<td>0.818</td>
<td>(0.647 - 1.034)</td>
</tr>
<tr>
<td>Gender</td>
<td>0.2103</td>
<td>1.668</td>
<td>(0.749 - 3.714)</td>
</tr>
<tr>
<td>Race</td>
<td>0.6108</td>
<td>1.314</td>
<td>(0.459 - 3.757)</td>
</tr>
<tr>
<td>Language</td>
<td>0.4188</td>
<td>0.695</td>
<td>(0.288 - 1.679)</td>
</tr>
<tr>
<td>Pn_meals_wk***</td>
<td>0.0144*</td>
<td>0.350</td>
<td>(0.151 - 0.811)</td>
</tr>
</tbody>
</table>

*Significant at p

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**Conclusion**

A statistically significant (p<0.0144) relationship exists between parental meal planning habits and elevated BMI in their children. Parents who disagreed with taking time to plan meals were 0.35 times as likely to have overweight children than parents who did take time to plan meals, with a confidence interval of 0.151-0.811.

**Conclusions** It appears that there is a correlation between a non-habit of parental meal planning and childhood obesity. This outcome is reasonable because parents who do not plan meals ahead of time may not shop for groceries beforehand or thoughtfully address healthy food choices containing necessary nutrients. Further exploration is needed to determine exactly what aspect of not taking time to plan meals earlier is correlated to increased children’s BMI.

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**Purpose of study** According to a 2013 report, the number of Oregon children on Medicaid who received an annual wellness exam by their pediatrician was 41%. We suspect that the number of adolescents who visit their primary care provider annually may be lower in rural Oregon for students who are on an Individualized Education Plan (IEP, indicating some form of learning disability). Previously published literature specifically linking students on an IEP and annual wellness exams was not found. The purpose of our project was to determine what barriers, if any, exist to healthcare for students on an IEP in rural Oregon.

**Methods used** Parents of students ages 14-18 who qualified for an IEP per district guidelines were given an 8-question survey to complete during an annual IEP meeting. Results were analyzed with a Chi-Square test after all surveys had been completed.

**Summary of results** 46 out of 140 parents filled out the survey (33%). 82% of respondents indicated that they had seen a provider in the past year. There was no significant difference in male vs. female students (p=0.6131) or in age groups (p=0.9761). 40% responded that they had sought mental health resources for their child. Of those that had sought mental health resources, 77.7% were able to see a mental health provider. There was no significant difference between male vs. female in seeking mental health resources (p=0.3293) or being able to see a provider (p=0.2521).

**Conclusions** We found that IEP students in this rural Oregon population were receiving annual wellness checks at a higher rate than the state Medicaid average of 41%. We suspect that our study had significant selection and non-response bias. Further study is needed to determine if these findings are generalizable.
Purpose of study
Beginning with the winter 2010-11 season, we implemented a care process model (CPM) to encourage hospital discharge with oxygen therapy for patients with bronchiolitis. Patients with diagnosis codes for chronic lung disease, tracheostomy dependence, or pulmonary hypertension were excluded, as were encounters that occurred outside of winter respiratory season (May-October). The primary outcome was mean length of stay. Balancing measures included number of hospital readmissions per episode of care, duration of hospital discharge, and diabetes (DM) as a known risk factor for readmission. In this project we assessed the impact of a home oxygen therapy CPM over a 10-year period.

Methods used
We conducted a retrospective cohort study of first year after implementation. We evaluated the impact of a home oxygen therapy CPM on hospital length of stay and cost of hospitalization in the year after implementation. A previous study suggested initial reductions in length of stay and cost of hospitalization in the first year after implementation. A total of 9,887 patients met inclusion criteria (3,033 pre-implementation; 6,854 post-implementation). Implementation of a home oxygen therapy CPM was associated with immediate decreases in mean length of stay (difference -11.1 hours, 95% CI -12.8 to -9.5 hours) and mean cost per episode of care (difference -$768, a 7.8% decrease, 95% CI -$1,262 to -$273), without an immediate change in readmission rate (difference 0.2%, 95% CI -0.3 to 0.7%). These findings have been sustained for 7 years after implementation and represent more than $5 million in hospital cost savings.

Summary of results
The home oxygen therapy CPM has demonstrated sustained decreases in hospital length of stay and cost without an increase in readmissions.

Conclusions
Clinicians value longer duration of follow-up, nursing and social work support, and information about medications and early intervention services. Future work will compare results with those of patients’ families.

Purpose of study
Discharge from a neonatal intensive care unit (NICU) requires significant medical and social support. Discrete choice experiments (DCE) are based in economic theory and are used to estimate an individual’s preferences when faced with a choice scenario. We used a DCE to identify key features of a NICU transition-of-care (TOC) program among NICU clinicians nationally.

Methods used
We designed a DCE survey containing 6 attributes of various levels based on qualitative focus groups with families of NICU infants: follow-up duration (3-24 months), financial support ($500-$4000/month), medical communication (appointment reminders, printed care plan, or directory of specialists), resource information (nutrition, housing, transportation, and financial aid), social support (peer, nursing, social work, or mental health), and medical information (durable medical equipment, diagnoses, medications, or early intervention services). Clinicians chose a preferred TOC program from 2 alternatives composed of varying levels of each attribute. Responses were analyzed via mixed logit models, and odds-ratios and willingness-to-pay for each attribute were estimated.

Summary of results
234 clinicians completed the survey. Each month of follow-up was worth $113. Appointment reminders were preferred vs. care plans or directory of specialists (OR=1.03, p=0.08 and 2.33, p<0.0001 respectively). Compared to housing, transportation, and financial aid information, nutrition information was more preferred (OR=1.35, 1.39, and 1.92 respectively, p<0.05). Nursing and social work support were preferred over peer support (OR=1.24 and 1.43, p<0.05) and worth $2,368 and $3,911, but mental health support was not (OR=0.93, p=0.06). Compared to durable medical equipment information, medication and early intervention services information were preferred (OR=1.19 and 2.08 respectively, p<0.05) and worth $1855 and $7992 while diagnosis information was not (OR=0.68, p=0.01).

Conclusions
Clinicians value longer duration of follow-up, nursing and social work support, and information about medications and early intervention services. Future work will compare results with those of patients’ families.

Purpose of study
Readmissions to the hospital are considered an indicator of quality of care. Pneumonia is a common cause of hospitalization, and diabetes (DM) is a known risk factor for both admission and readmission in patients with pneumonia. However, there is a lack of studies specifically evaluating readmission risk factors and fatal readmission risk factors in patients with DM originally admitted for pneumonia. Determining these risk factors may direct efforts in decreasing readmission for these common diagnoses. In this project we determined the risk factors for readmission in patients with pneumonia and DM.

Methods used
The 2013 National Readmission Database (NRD) was used to identify DM adults discharged alive after hospitalization for pneumonia (weighted N=229849), whether these adults were readmitted within 30 days of discharge, and mortality during the readmission. We identified readmission risk factors including comorbidities, procedures, and...
demographics. We also determined risk factors specifically for readmissions resulting in mortality.

Summary of results Overall, 18.1% of DM patients were readmitted. Top three readmission reasons were pneumonia (14.7%), heart failure (9.4%), and septicemia (8.7%). This differed from non-DM patients in which the top three readmission reasons were pneumonia (17.2%), septicemia (9.2%), and COPD (6.5%). In DM patients, independent risk factors for readmission included metastatic cancer (adjusted OR 1.68), solid tumor (aOR 1.61), drug abuse (aOR 1.56), and hemodialysis (aOR 1.38). Independent protective factors included private insurance (aOR 0.78). Independent risk factors for mortality during readmission included metastatic cancer (aOR 3.62), solid tumor (aOR 2.36), age >65 (aOR 2.03), blood transfusion (aOR 1.96), lymphoma (aOR 1.72), and weight loss (aOR 1.65).

Conclusions Our study demonstrates that pneumonia is the leading reason for readmission in both DM and non-DM patients. In addition, patients with DM are at higher risk for readmission due to heart failure compared to non-DM. The risk factors could be used to identify patients who are at higher risk such as focused specific post discharge support. Since drug abuse is a modifiable risk factor, interventions to decrease drug abuse in DM patients may decrease readmission rates.

TIMING OF ADDICTION CONSULTATION IN HOSPITALIZED PATIENTS STARTING BUPRENORPHINE

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Purpose of study A growing body of evidence supports initiation of medications for OUD, including buprenorphine, during hospitalization. It is unclear what factors during hospitalization are associated with likelihood of linkage with ongoing addiction treatment after discharge. This study explores the associations between early addiction consultation and medication initiation during hospitalization and successful linkage with outpatient treatment.

Methods used A retrospective chart review was completed at an urban, academic hospital for hospitalized patients with opioid use disorder. All hospitalized patients with opioid use disorder who were evaluated by the addiction medicine consult service from January 2018 to June 2019 and agreed to start buprenorphine were included. Timing of consultation was determined based on the proportion of hospitalization completed (days from admission to consult ÷ length of stay). Binary logistic regression was used to compare timing of consultation with inpatient buprenorphine initiation. An additional logistic regression was used to compare the odds of successful linkage (defined as any outpatient follow up within 30 days of discharge) between patients initiated on buprenorphine during hospitalization with those receiving a prescription at discharge. All analyses were completed with Stata, version 16.

Summary of results During the study period, 145 patients met eligibility criteria. Individuals with earlier addiction consultation were more likely to start buprenorphine during hospitalization compared to those evaluated closer to discharge (p=0.000, OR=0.04). There was no significant difference in successful linkage between the 115 patients with inpatient buprenorphine induction versus 30 patients that started at discharge (p=0.15, OR=1.89).

Conclusions Hospitalized patients with OUD who have earlier addiction consultation are more likely to have inpatient buprenorphine induction. Additional engagement with clinicians and opportunities to discuss treatment goals may allow patients to start buprenorphine early. Inpatient initiation of buprenorphine is not associated with increased odds of linkage within 30 days of discharge compared to providing a prescription of buprenorphine at discharge.

DEEP LEARNING CLASSIFICATION OF PROSTATE MRI SEQUENCES

1PB Bhattar*, 2M Bards, 2C Chahine, 2A Ushinsky, 2D Fujimoto, 2WA Grant, 1P Chang, 1R Houshyar. 1University of California Irvine Medical Center, Orange, CA; 2Washington University, St. Louis, MO

Purpose of study An early imaging preprocessing step involves uniformly labeling data sources. For MRIs, this represents a...
challenge due to the (1) heterogeneity in vendors, sequences, and protocols, and (2) potential mislabeling that has been reported to be as high as 10%. Furthermore, for multiparametric assessment for prostate MRIs, this can be a hurdle for research and clinical studies that rely on comparisons. Therefore, we develop a neural network that can automatically label sequences.

Methods used This retrospective study included patients who had a prostate MRI and an ultrasound MRI fusion transrectal biopsy between September 2014 and August 2018. For this study, the DICOM header information was used to classify 57,063 images as either T2W, DWI, or DCE. The number of T2W, DWI, and DCE images used were 17,195, 23,576, and 5,918 respectively. The convolutional neural network (CNN) used 46,689 images for training and 11,412 images for validation. The two-layered convolutional neural network was then trained for 8,000 iterations (figure 1). A fully connected layer with one hidden layer was used before the output layer. The softmax cross entropy loss function was used to adjust the weights during training.

Summary of results In validation, the CNN accurately classified 11,410 out of 11,412 images for an accuracy of 99.2%.

Conclusions A CNN can be accurately used to identify the sequences of T2W, DWI, or DCE. By using a CNN to complete the series identification, the process of automatically examining a prostate mpMRI and determining the PI-RADS score for a lesion will be expedited. The time needed to classify the series will be reduced and the DICOM header descriptions that are entered by the technologists can be ignored.

Purpose of study Cranial computed tomography uncovers a variety of injuries in pediatric blunt head trauma patients. However, cranial CT poses many risks for these children and compliance with the Pediatric Emergency Care Applied Research Network (PECARN) recommendations seeks to reduce the number of CT scans ordered without missing significant traumatic brain injuries. The aim of this project is to assess the degree of compliance with the PECARN criteria following blunt head trauma among children in a single emergency department and identify the common findings that prompted cranial CT utilization over observation.

Methods used This is an observational cross-sectional study looking at patients under the age of 18 who received cranial CT scans over 6 months in a single county emergency department. Via retrospective chart review, we determined whether or not PECARN criteria were met and identified symptoms common among patients in the category in which either CT or observation would be acceptable.

Summary of results 253 head CTs were obtained in the pediatric emergency department from January to June of 2019—68 without head trauma, 24 with unknown head trauma, and 137 with recorded head trauma. Among the 137 patients with head trauma, 69.3% of the patients were categorized as ‘CT vs. observation’ under PECARN criteria, 21.2% as PECARN negative, and 10.2% as PECARN positive. 7 out of 95 CT vs. observation patients were discovered to have traumatic brain injuries such as subdural hemorrhages and fractures on CT, although none required neurosurgical intervention. Among these 95 patients, we identified the most common presenting symptoms of headaches, loss of consciousness, vomiting, nausea, and lethargy. Headaches were reported in 55% of the CT vs. observation cases, loss of consciousness in 40%, and vomiting in 26%.

Conclusions In accordance with PECARN criteria, cranial CT may be over-utilized in pediatric head trauma patients presenting to this hospital site, as there were a number of patients who did not meet PECARN criteria but had a cranial CT performed. We will use quality improvement methodology to decrease the utilization of cranial CT in those who do not meet PECARN criteria. We will additionally look to discover what drives decision-making processes for providers when a patient is in the CT vs. observation category.

Purpose of study With an increase in sport-related concussions and heightened accessibility of technology, computer-based assessments via direct entry have become widely used among sports medicine professionals. The advantages of direct participant entry via computer over person to person interview protocols include instantaneous results and a greater capacity to gain larger study populations via ease of response entry and removal of the need for a qualified interviewer. In addition, privacy among participants via the direct computer entry may limit some bias as a consequence of threat of disclosure and social desirability related to in person interviews. Currently no studies exist which compare the validity of results of SCAT using computer-based vs in person interview, thus the aim of this study was to compare the differences between collection methods.

Methods used Three hundred and eighty-five (N=385) National Collegiate Athletic Association Division III student-athletes from CLU completed the SCAT symptoms via face to face interview thus the aim of this study was to compare the differences between collection methods.
females ranged from 0.091 – 0.529 (P<0.001), demonstrating poor-to-moderate strength.

Conclusions Overall, these results demonstrate a significant relationship between direct computer entry and face to face delivery of the SCAT for both group and gender. In addition, these findings suggest that the direct computer entry platform is as accurate as the in-person interviews, and either method of SCAT administration would be reliable for use by medical practitioners in determining athlete concussion symptoms and status.

THE ROLE OF DO NOT ATTEMPT RESUSCITATION

Abstracts

THE ROLE OF DO NOT ATTEMPT RESUSCITATION

1R Winchell, 1J Blackwood, 1CR Counts, 2D Drucker, 1A Jennerich, 1K Pompeo, 1J Waldron, 5S Feder, 1T Rea, 1M Eisenberg, 1M Sayre, 1P Kudenchuk. 1University of Washington, Seattle, WA; 2Seattle and King County, Division of Emergency Medical Services, Seattle WA; 3Shoreline Fire Department, Shoreline, WA; 4King County Medic One, Seattle, WA; 5King County EMS, Seattle, WA

Purpose of study A Do Not Attempt Resuscitation (DNAR) directive is a physician order that conveys patient wishes to not be resuscitated from a cardiac arrest. Emergency medical services (EMS) providers encounter DNAR orders when managing out-of-hospital cardiac arrest (OHCA). Little is known about their prevalence or how they might affect care provided.

Methods used We retrospectively analyzed all treated adult OHCA patients in 2018 that had DNAR orders per EMS in a large urban community. Data were abstracted from EMS PCRs, 911 dispatch audio recordings, and defibrillator ECG and audio recordings. We compared patients who received full resuscitative efforts to those with some limitation of their resuscitation.

Summary of results Of the 98 included patients, the average age was 80 years (SD ±10.6), 54 (55%) were male, 54 (55%) had their OHCA at home, and 53 (54%) received bystander CPR. Twenty-three patients (23.5%) received full efforts, while 75 (76.5%) received limited efforts. The full efforts group were more likely to have an initially shockable rhythm (17% vs. 7%), and twice as likely to have a witnessed arrest (70% vs. 33%, p<0.05). Twenty patients (87%) who received a full resuscitation had limited or no interventions on their DNAR.

Conclusions Delays in the presentation of DNAR orders during OHCA are common. This delay combined with family member confusion/contradiction of the DNAR orders results in undesired resuscitation. How DNAR directives are best operationalized and implemented requires further evaluation.

COEXISTING CHRONIC DISEASE INCREASES THE RISK OF MISCLASSIFICATION OF TYPE 1 DIABETES

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Purpose of study An accurate diagnosis of diabetes type in the Electronic Medical Record (EMR) is essential for effective treatment, prevention of medical errors, and the validity of research that depends upon these large databases. We hypothesize that Type 1 Diabetes (T1D) patients with chronic comorbidities will be more likely to be misclassified as having Type 2 Diabetes (T2D) compared to those without such comorbidities.

Methods used Data were collected from HealthFacts® from 2006 to 2017 from patients diagnosed with T1D or T2D using ICD-9-10 codes. Of those, 9,676 patients were coded for T1D alone and were compared to 271,428 patients who were coded for both T1D and T2D. Patients coded for T2D alone were excluded from analysis. Multiple logistic regression was used to estimate the odds of misclassification for patients with coexisting comorbidities, including hypertension, cardiovascular disease (CVD), asthma, chronic obstructive pulmonary disease (COPD), chronic kidney disease (CKD), alcohol use disorder (AUD), and depression as compared to those without such comorbidities. The model is adjusted for age, sex, race, urban place of residence, marital status, and health insurance.

Summary of results Of 3.3 million diabetes patients, 8.3% were misclassified at least once. Of 281,104 T1D patients, 47.9% had coexistent hypertension, 54.4% had CVD, 1.1% had asthma, 13.4% had CKD, 6.7% had COPD, 9.0% had depression and 1.8% had AUD. In general, patients with chronic disease were at higher risk for misclassification of diabetes (see table 1). For example, patients with a history of CVD or depression had significant 2.24-fold and 2.20-fold increases, respectively, in the odds of diabetes misclassification compared to those without such a history.

<table>
<thead>
<tr>
<th>Comorbidity</th>
<th>Adjusted OR for Misclassification of T1D (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>CVD</td>
<td>2.24 (1.98-2.54)</td>
</tr>
<tr>
<td>Depression</td>
<td>2.20 (1.96-2.46)</td>
</tr>
<tr>
<td>CVD</td>
<td>1.94 (1.74-2.15)</td>
</tr>
<tr>
<td>AUD</td>
<td>1.65 (1.31-2.08)</td>
</tr>
<tr>
<td>COPD</td>
<td>1.50 (1.29-1.74)</td>
</tr>
<tr>
<td>Hypertension</td>
<td>1.25 (1.09-1.43)</td>
</tr>
<tr>
<td>Asthma</td>
<td>1.13 (1.09-1.30)</td>
</tr>
</tbody>
</table>

Conclusions A history of comorbid hypertension, CVD, COPD, CKD, AUD, and/or depression significantly increases the risk of misclassification of T1D.
Hematology and oncology II
Concurrent session
10:15 AM
Friday, January 24, 2020

316 FUSION-NEGATIVE ONCOGENES RESCUE FUSION-POSITIVE Rhabdomyosarcoma from GATOR2 dependence

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10.1136/jim-2019-WMRC.316

Purpose of study Rhabdomyosarcoma (RMS), the most common pediatric soft-tissue sarcoma, is comprised of two genetic subtypes. Fusion-positive (FP) RMS is characterized by PAX3-FOXO1 or related translocations, while fusion-negative (FN) RMS harbors mutations in the RAS/P13K pathway. Patients with high-risk RMS have poor outcomes. Characterizing the molecular connections between genomic drivers and cellular fitness may enable future therapeutic advancements in RMS.

Using a CRISPRi screen, we identified two components of the GATOR2 signaling complex, an activator of mTOR, as necessary in FP RMS but not FN RMS. We hypothesize that FN oncogenes are sufficient to overcome dependence on GATOR2 for mTOR-regulated proliferation.

Methods used To characterize the effects of FN oncogenes on GATOR2 loss, we conducted competition growth assays using FP RMS cells, and FP cells transduced with the FN oncogenes NRASQ61H and FGFR4V550E. We used CRISPRi to knockdown GATOR2 components in these cells and compared the growth of parental and oncogene-modified cell lines. We measured mTOR activation in these conditions by immunoblotting for p70S6K phosphorylation and 4EBP1 binding to m7-GTP sepharose beads.

Summary of results We confirmed that knockdown of GATOR2 components in FP RMS cells decreased their growth compared to control cells. However, introduction of FN oncogenes partially rescued cells from GATOR2 loss. This was also seen with inactivation of the negative mTOR regulator TSC1. Immunoblots showed that FN oncogenes did not rescue p70S6K phosphorylation after GATOR2 loss. By contrast, they did induce dissociation of 4EBP1 from m7-GTP, suggesting a restoration of cap-dependent translation.

Conclusions Our data demonstrate that loss of GATOR2 in FP RMS limits mTOR activation by environmental amino acid signaling. However, we find that FN oncogenes are sufficient to rescue cells from GATOR2 loss. Additionally, our data indicate that reduced cap-binding by 4EBP1, but not phosphorylation of p70S6K, is tied to the survival and proliferation of FP RMS. Based on our findings, we plan future investigation of the specific roles of 4EBP1 in RMS. Our work provides molecular insights into the pathways upstream (eg, GATOR2) and downstream (eg, 4EBP1) of mTOR which are critical for RMS survival.

317 EXPLORING T2-FLAIR MISMATCH AMONG IDH-MUTANT ASTROCYTOMAS: PATTERNS OF EVOLUTION AND FURTHER CHARACTERIZATION

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10.1136/jim-2019-WMRC.317

Purpose of study The T2-FLAIR mismatch sign (T2FM) is a newly defined biomarker on MRI which demonstrates high predictive value for isocitrate dehydrogenase (IDH) mutant astrocytomas among lower grade gliomas, helping facilitate early diagnosis and treatment planning. The purpose of this study is to further explore T2FM by characterizing tumors which display the sign, evaluating the predictive value of its various definitive criteria (T2 homogeneity and hyperintense FLAIR rim), and describing patterns in its temporal development.

Methods used Medical records from 64 patients with astrocytomas treated at the Alvord Brain Tumor Center were assessed for age at diagnosis, gender, WHO grade, molecular status, seizure history, tumor characteristics on pre-treatment CT, MRI, and pathology, documentation of T2FM, treatment course, and temporal changes in tumor appearance. Cases were divided into those meeting ‘classic’ T2FM definitive criteria (homogenous T2, FLAIR rim positive), those considered ‘ambiguous’ and not fulfilling all criteria (heterogeneous T2, FLAIR rim positive), and those which were T2FM negative (FLAIR rim negative). Groups were compared using Chi square and qualitative analyses.

Summary of results Including ‘ambiguous’ tumors increased T2FM sensitivity for IDH mutant astrocytomas 30% without decreased specificity. Tumors with T2FM characteristics were more cystic, less enhancing, and affected younger patients. T2FM was 100% persistent in subtotally resected residual tumors and disappeared with radiotherapy, persisted in 5/8 recurrent tumors which were originally T2FM positive, and was identified in tumors with high grade characteristics. Trends were identified among patient age, tumor location, mismatch status, and IDH status.

Conclusions A hyperintense FLAIR rim, regardless of T2 appearance, demonstrates high predictive power for IDH mutation among astrocytomas and may be used to determine T2FM status. Additionally, T2FM evolves in predictable patterns and may be seen in higher grade gliomas, findings which have not been previously described. Lastly, trends among patient age, tumor location, and T2FM may allow inference of IDH status when sequencing information is not available, further expanding its clinical utility.

318 EPHRINB2 AND EPHB4 MODULATES INVASION AND PROLIFERATION PATHWAYS IN Glioblastoma Multiforme

S Bhatia, S Bukkapatnam*, S Karam. University of Colorado School of Medicine, Aurora, CO
10.1136/jim-2019-WMRC.318

Purpose of study The aggressive nature of Glioblastoma Multiforme may be attributed to the complex invasive and proliferative pathways that these tumors commandeer to promote tumorigenesis. EphrinB2, a receptor tyrosine kinase, has
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319 CYTOKINE AND GROWTH FACTOR PROFILING FOR THE DIFFERENTIATION OF MULTIPLE MYELOMA PATIENTS FROM THOSE WITH MONOCLONAL GAMMOPATHY OF UNKNOWN SIGNIFICANCE

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Purpose of study To determine if we could differentiate individuals with multiple myeloma (MM) from those with monoclonal gammopathy of unknown significance (MGUS) based on bone marrow (BM) and peripheral blood (PB) plasma cytokine and growth factor profiles. MM is a disease of dysplastic plasma cells that localize to bone marrow, producing high levels of immunoglobulin heavy and light chains, which can lead to hypercalcemia, renal disease, anemia, and bone lesions. MM is often preceded by increased immunoglobulin production but no clinical symptoms referred to as MGUS. IL-16 is an important growth factor in the development of myeloma with B cell activating factor (BAFF), and IL-17 related to the increase in IL-16.

Methods used A multiplexed assay was developed for IL-2r, IL-6, IL-8, IL-16, IL-17, VISTA (V-domain Ig suppressor of T cell activation) and BAFF. Bone marrow and plasma samples obtained from 154 diagnosed multiple myeloma and 16 MGUS patients were tested.

Summary of results BAFF expression was significantly elevated in MM compared to MGUS patients in both BM (3,529 vs 1,430 pg/mL, p<0.05) and PB (2,167 vs 948 pg/mL, p<0.05). IL-16 and IL-2r were also more strongly expressed in MM vs MGUS, but statistical differences were only observed in the BM samples. IL-16 BM (378 vs 124 pg/mL, p<0.05) and IL-2r BM (1,001 vs 124 pg/mL, p<0.05). Matched BM and PB samples from the MM patients demonstrated no strong correlation for BAFF (Slope 0.349, r² 0.362), IL-16 (Slope 0.014, r² 0.029) or IL-2r (Slope 0.313, r² 0.244).

Conclusions IL-16 and BAFF, known tumor promoting factors, and IL-2r, an immune inflammatory marker, were significantly increased in multiple myeloma when compared to MGUS patients. All markers investigated were expressed in greater concentrations in the BM than the peripheral blood, but no direct correlation was observed. Monitoring IL-16, BAFF and IL-2r marker profiles of MGUS patients may allow us to predict the onset of Multiple Myeloma and may potentially lead to anti-cytokine therapy for this malignancy.

320 THE EFFECT OF PREVIOUS CHEMOTHERAPY TREATMENT ON OVARIAN FOLLICLE COUNT IN OVARIAN TISSUE CRYOPRESERVED FOR FERTILITY PRESERVATION

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Purpose of study In the last several years, ovarian tissue cryopreservation (OTC) has become a more widely accepted method of fertility preservation for children and young adults who are at high risk of infertility as a result of cancer treatment but who are not able to store mature eggs. In some fertility preservation programs, prior exposure to chemotherapy is an absolute exclusion criterion. In the Oxford Programme, patients are eligible for tissue storage if they have undergone chemotherapy prior to collection. This abstract presents the result of comparative analysis of follicle counts across this cohort of patients.

Methods used The pathology data from 306 female patients enrolled in the Oxford Reproductive Tissue Cryopreservation Programme (ORTC) between September 2013 to June 2019 was analyzed. This cohort included patients between the ages of 2 months to 38 years old who fulfilled the ORTC eligibility criteria. Follicle counts were analyzed by age, prior exposure to chemotherapy and diagnosis.

Summary of results All patients regardless of age, diagnosis or previous exposure to chemotherapy had follicles within the tissue analyzed in the pathology laboratory. Age, across all data sets, had the most significant effect on follicle numbers. For patients who had previously been exposed to chemotherapy, while the follicle count across the age range was lower than age matched patients who had not had pre-chemotherapy treatment, the effect of previous chemotherapy was most significant in the older patient cohort. Diagnosis as an independent variable did affect follicle numbers per se.

Conclusions These results demonstrate that ovarian tissue collected from patients both pre- and post-chemotherapy contains significant numbers of primordial follicles. Age had a more
significant effect on follicle numbers than previous exposure to chemotherapy. This data supports the inclusion of patients who have had non sterilizing treatment in ovarian tissue cryopreservation programs for children and young adults.

**Abstract Withdrawn**

**USE OF AROMATHERAPY AS A COMPLEMENTARY THERAPY IN ALLEVIATING CHEMOTHERAPY RELATED ADVERSE EFFECTS IN CANCER PATIENTS**

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10.1136/jim-2019-WMRC.322

Purpose of study Use of aromatherapy in relieving anxiety and pain in different settings has been reported but its use in alleviating the side effects related to chemotherapy or radiation therapy in cancer patients is less clear. The purpose of this study was to investigate the effectiveness of aromatherapy in alleviating the adverse effects related to chemotherapy or procedures in cancer patients.

Methods used A literature review using online resources including Pubmed and Google Scholar with the keywords ‘cancer aromatherapy’ was performed. Only studies of adult cancer patients with a control group published after 2005 were included in our analysis.

**Summary of results** Of the 20 articles, only 7 met our inclusion criteria (table 1 below). The primary reason for exclusion was the lack of a control group. The types of essential oils and length of aromatherapy treatment was variable among the studies. In 5/7 studies, aromatherapy showed improved in symptoms, such as nausea, vomiting or pain but the beneficial effect on vital signs was less clear. In addition to the included studies, we searched for reports of side effects associated with aromatherapy. Allergic reactions were seen in minority of patients, and one study found an association between nosocomial outbreak of drug resistant *Pseudomonas* infections and use of inhalation aromatherapy.

Conclusions Our review suggests aromatherapy may have a role in alleviating some of the symptoms associated with chemotherapy/radiation therapy in adult cancer patients. However, due to potential serious side effects, caution should be taken in the use of aromatherapy in cancer patients. Larger studies that control for the type of aromatherapy and other confounding variables are needed.

**Abstract 322 Table 1** Summary of studies on use of aromatherapy in cancer patients

<table>
<thead>
<tr>
<th>1st Author; Year</th>
<th>Variable Studied</th>
<th>Aromatherapy (AR) Used</th>
<th>Patient’s Type of Cancer</th>
<th>Length/Frequency of Treatment</th>
<th>Number of Control and Number of Aromatherapy (AR) Subjects</th>
<th>Results</th>
</tr>
</thead>
<tbody>
<tr>
<td>Heydarkarad; 2019</td>
<td>Quality of Sleep</td>
<td>Rosa Damascena (Inhalation)</td>
<td>Any Type of Cancer</td>
<td>2 weeks; AR treatment nightly</td>
<td>18 control 18 oil 5% 18 oil 10%</td>
<td>Duration and quality of sleep better in AR groups*</td>
</tr>
<tr>
<td>Nakayama; 2016</td>
<td>Salivary gland damage measured by scintigraphy</td>
<td>1.0 mL Citrus Limon and 0.5 mL Zingiber Officinal (Inhalation)</td>
<td>Thyroid Cancer</td>
<td>2 weeks inhalation; AR treatment 10 minutes before each meal</td>
<td>36 control 35 AR group</td>
<td>Higher rate of accumulation/secretion in parotid and submandibular glands in AR*</td>
</tr>
<tr>
<td>Iter; 2019</td>
<td>Vital signs, pain during port insertion</td>
<td>Diluted orange, chamomile, and lavender oil in 70 ml water (Inhalation)</td>
<td>Any Type of Cancer</td>
<td>AR inhalation treatment during procedure</td>
<td>30 control 30 AR group</td>
<td>* Significantly decreased pain experienced during the procedure* - Effect on vital signs not statistically significant</td>
</tr>
<tr>
<td>Tamaki; 2018</td>
<td>Quality of life, vital signs, &amp; sleep quality</td>
<td>Aroma oils including ylang-ylang, orange, &amp; lavender (Inhalation)</td>
<td>Breast Cancer</td>
<td>Aroma oil placed for inhalation for 9 hours prior to surgery</td>
<td>52 control 110 AR group</td>
<td>Results showed no effects of aromatherapy on quality of life, vital signs, or sleep quality</td>
</tr>
<tr>
<td>Wilkinson; 2007</td>
<td>Anxiety &amp; depression</td>
<td>Choice of 20 essential oils (Topical - massage)</td>
<td>Any Type of Cancer</td>
<td>4 weeks; 1 hour weekly sessions of AR massage</td>
<td>144 control (usual care alone) 144 supportive care + AR massage</td>
<td>AR improves anxiety/depression for 2 weeks* but had no long-term benefits</td>
</tr>
<tr>
<td>Zorba; 2018</td>
<td>Nausea &amp; vomiting</td>
<td>Essential oil mixture of peppermint, bergamot, and cardamom in almond carrier oil (Inhalation &amp; Topical - massage)</td>
<td>Breast cancer</td>
<td>20 minute AR foot massage (massage group) &amp; 3 minute inhalation aromatherapy (inhalation group) before each patient’s chemotherapy cycles</td>
<td>25 control 25 massage AR 25 inhalation AR</td>
<td>Both inhalation and massage aromatherapy treatment before chemotherapy cycles showed to significantly decrease nausea compared to control*</td>
</tr>
<tr>
<td>Rostami; 2019</td>
<td>Peripheral Nephropathy</td>
<td>Topical Citrullus extract oil (Massage)</td>
<td>Breast Cancer</td>
<td>2 months; 2 massages to hands and feet per day</td>
<td>15 control 17 AR group</td>
<td>No significant difference in placebo and AR group</td>
</tr>
</tbody>
</table>

* p<0.05
Purpose of study Hepatocellular carcinoma (HCC) is the fourth-leading cause of cancer-associated death, claiming 800,000 lives each year globally. At the same time, an increasing number of patients are using the Internet as a source of health information. However, limited research has been done on assessing the quality of HCC websites. Therefore, we aim to systematically evaluate the quality of online HCC information.

Methods used The term ‘hepatocellular carcinoma’ was searched using the search engine Google and the meta-search engines Dogpile and Yippy. The overall highest-ranked 100 websites were extracted based on pre-specified inclusion and exclusion criteria. A previously validated tool was used to evaluate quality based on several parameters including affiliation, accountability, interactivity, structure & organization, readability, and content quality.

Summary of results The search yielded over 1,100 websites. Of the top 100, 53% were commercial. Although 95% disclosed ownership and sponsorship, other measures of accountability were poor – only 30% identified their author(s), 42% cited sources, and 33% were updated within the past two years. Average readability was judged to be at a grade 11.8 level using the Flesch-Kincaid grading system, which is significantly higher than the recommended grade-six level (p<0.0001). Definition and symptoms were the most commonly covered topics, while prognosis and prevention were the least covered. Content accuracy was high among covered topics, except for incidence, for which 23% had mostly inaccurate information. Overall, non-commercial websites were higher in quality compared to commercial ones (p<0.002).

Conclusions Although website contents are generally accurate, authorship, attribution, currency, and coverage are currently deficient among many HCC websites. Additionally, difficult readability may pose a barrier for patient comprehension. Despite high public demand for information regarding prevention and prognosis, very few websites cover those topics. Healthcare professionals should be aware of the limitations of online HCC information, so they can be proactive in guiding patients to reliable resources to address their specific needs.

A RARE CASE OF ABDOMINAL PAIN:RELATED TO IDIOPATHIC LEFT OVARIAN VEIN THROMBOSIS

1GS Petersen, 1M Kalkur, 1N Trang*, 1Kem Medical, Bakersfield, CA; 2UCLA, Los Angeles, CA

Purpose of study Ovarian vein thrombosis (OVT) first described in 1956 is a rare thrombotic condition with an incidence of 60-fold lower compared with leg DVT, yet could be potentially life threatening. Upon literature review, OVT cases are most often seen on the right ovarian vein. We describe an unusual case of idiopathic OVT, probably the first reported case of idiopathic left OVT.

Methods used Retrospective case report

Summary of results We report the case of a 47 year old G7P7 Caucasian female with no past medical history who presented to a local hospital for acute onset of back pain radiating to her ribs and colicky left lower quadrant pain for 1 day. She denied any associated symptoms. CT scan of the abdomen and pelvis with IV contrast revealed left gonadal vein thrombosis. She was then transferred to our facility for further management and treatment. On admission, patient was afebrile and hemodynamically stable. Pelvic ultrasound showed normal endometrium, no mass or cyst in the uterus or ovary with left ovary measuring 2.4cm x 1.5cm with normal blood flow. Work up for inherited and acquired thrombophilia was obtained and later proved unremarkable. Therefore, the diagnosis of OVT was made and anticoagulation treatment was initiated with low molecular weight heparin (Levonox 1mg/kg q12h). Patient’s symptoms completely resolved in the next 24 hours and she was discharged home on direct oral anticoagulant (Rivaroxaban) for 3 months. At her two month follow-up appointment, CT of abdomen and pelvis with IV contrast showed complete resolution of the left ovarian vein thrombosis.

Conclusions OVT is an uncommon yet potentially life threatening thrombotic condition with a 25% risk of developing into PE and 4% mortality rate. Based on our research, 70-90% OVT cases occur on the right ovarian vein, whereas 11-14% are bilateral. OVT affects mostly postpartum women but might also be associated with pelvic inflammatory disease, malignancies, and pelvic surgical procedures. This is probably the first case reported for idiopathic left OVT in a healthy woman. Standard guidelines for managing OVT are lacking. Historically, heparin followed by bridging to warfarin until a therapeutic INR is achieved was the main stay of treatment. Recently, direct oral anticoagulants offers a potentially useful alternative even the efficacy of these drugs has not been studied yet.

Infectious diseases II

Concurrent session

10:15 AM

Friday, January 24, 2020

MEASLES VACCINE RESPONSE VARIES IN 12-MONTH-OLD KENYAN CHILDREN WHICH MAY INCREASE MEASLES TRANSMISSION

DS Mason*, R Rodford, University of Colorado School of Medicine, Aurora, CO

Purpose of study Measles is a preventable infectious disease, but still presents a large burden of disease throughout the world. In 2017, measles caused roughly 110,000 deaths worldwide. The measles vaccine has greatly reduced measles-related morbidity and mortality. Despite this, Africa still suffers from this disease. Reduced measles vaccine efficacy has been reported in Sub-Saharan Africa. The aim of this study was to determine the response to measles vaccine following routine 9-month vaccination in infants living in a malaria endemic region of sub-Saharan Africa.

Methods used Plasma samples were collected from children from Kisumu, Kenya, at 12 months of age, three months after routine vaccination for measles. These plasma samples were tested by indirect ELISA to measure anti-measles antibodies.
Each sample was analyzed in triplicate. Samples that had one well with an optical density >0.5 different from the other two wells were trialed again to account for pipetting error. If after reanalysis the wells of the sample no longer had a well with a difference >0.5, the sample was included in analysis with its original wells excluding the well that differed >0.5. However, if the sample again had a well that differed in optical density from the other two wells by >0.5, the sample was considered an outlier and excluded from analysis. Eleven samples were excluded, leaving 143 samples included in the final analysis.

Summary of results The mean optical density was 1.83 (SD ± 0.72; 95% CI 1.71–1.94), with a minimum optical density of 0.42 and maximum optical density of 3.92. Twenty-two participants had an OD of 1.0 or less, raising concern of not being immunoprotective against measles.

Conclusions These twenty-two participants represent 15% of the total participant sample. For the measles vaccine to be protective, roughly 95% of the community must be vaccinated to prevent transmission. This then represents a possibility for transmission of measles to occur in these communities, particularly in young children who are more at risk of infection. The variability in levels of anti-measles antibodies following routine measles vaccination in this sample are not entirely understood and will be the subject of ongoing analysis.

326 INCORPORATING ZINC OXIDE NANOPARTICLES IN CONTACT LENS MATERIAL TO IMPROVE BIOCOMPATABILITY

1G Davis*, 1,2AG Gapeva, 1Cho, 2F Spons, 3Fuchs. 1Western University of Health Sciences, Pomona, CA; 2Christian Albrechts Universitat, Kiel, Germany; 3Western University of Health Sciences, Pomona, CA

10.1136/jim-2019-WMRC.326

Purpose of study Zinc oxide nanoparticles (ZON) have been found to filter UV radiation, prevent bacterial growth and encourage wound healing. Our purpose is to harness these useful properties by incorporating ZON into gels which are otherwise chemically identical to modern contact lenses. We will then test our enhanced contact lenses to see how their hydrophilicity, UV filtration and antimicrobial properties have changed. If successful, the implications may someday afford contact lens wearers a safer and more comfortable contact lens experience.

Methods used Gels chemically identical to modern contact lenses were produced with varying ratios of hydroxyethyl methacrylate (HEMA) and methacrylic acid (MAA), cross-linked with ethylene glycol dimethacrylate (EGDMA), and heat polymerized in presence of 4,4’-Azobis(4-cyanovalearic acid).

The wettability of these gels was measured by photographing a fully submerged, 2 cc, sessile air bubble as it contacted the gel surface in both deionized water and saline solutions. The angle between air bubble and gel quantified how varying ratios of HEMA:MAA impacted the hydrophilicity of the gels.

Summary of results To date, we have successfully produced polymerized material that resembles modern contact lenses in strength, flexibility, hydrophilicity and transparency. By experimenting with different ratios of HEMA:MAA and polymerization methods, we have established that a 70:30 HEMA:MAA ratio coupled with thermoinitiator consistently produces the strongest, clearest and most flexible lenses.

Conclusions Now that we have successfully established a method for polymerizing gels. We will be able to move to the next steps which include:

- Incorporating ZON into gels with the established 70:30 HEMA:MAA ratio.
- Comparing how ZON impacts lens wettability using the sessile drop technique.
- Growing bacteria on lenses to test antimicrobial properties of nanoparticles in lens polymer.

327 ASSESSMENT OF MUTANT STAPHYLOCOCCUS AUREUS AND MRSA STRAINS FOR DRUG LEAD OPTIMIZATION

1AC Lazarski*, 1N Molasky, 2J Gillespie, 3F Buckner. 1The University of Texas at El Paso, El Paso, TX; 2The University of Washington, Seattle, WA

10.1136/jim-2019-WMRC.327

Purpose of study Antibiotic resistance is an issue for the treatment of bacterial infections. S. aureus, especially methicillin resistant S. aureus (MRSA), has developed resistance to many current treatments. The increase in resistance demands a response to develop alternative therapies. The methionine t-RNA synthetase (MetRS) enzyme of bacteria is a novel drug target for many gram-positive species, including S. aureus. A series of small molecule inhibitors is being developed that bind this target. To understand and mitigate the risk of resistance developing to this new target, experiments were performed to characterize mutations that confer resistance.

Methods used Using tryptic soy agar plates containing four times the MIC of three different MetRS inhibitors, resistant S. aureus mutants were generated. Eight to 12 resistant colonies were selected and DNA sequenced to locate mutations in the MetRS gene. Mutations in the enzyme were modelled using I-TASSER and SWISS Model for visualization of the mechanism of resistance, and MICs of resistant strains were measured following CLSI guidelines.

Summary of results Mutant S. aureus colonies contained mutations I57N, V108M, H101P, and V105F in the MetRS gene. The magnitude of shifts in the MIC of inhibitors was dependent on the mutation present, with I57N causing the highest increase in MIC of the newly sequenced mutations. Molecular models of previously identified MRSA mutants elucidated potential mechanisms of resistance. Mutations affected one of two binding regions in the MetRS enzyme, the methionine binding pocket or auxiliary pocket (AP). Changes in the AP pocket caused greater shifts in MIC, but occurred less often. Assessment of the inhibitors MIC revealed that molecules with no halogen substituents in the AP binding region, or flexible regions, maintained better activity on mutant MRSA strains.

Conclusions To optimize MetRS inhibitors for S. aureus and other gram-positive bacteria, mechanisms of resistance and cross resistance were investigated. Through experiments on resistant strains, new MetRS inhibitors have been generated limiting the effects of resistance mutations. This work adds to our progress of developing a novel antibiotic class that is not susceptible to inducing antibiotic resistance.
Abstracts

328 PALMITIC ACID POTENTIATES MYCOBACTERIAL GROWTH IN MACROPHAGES
S Schwartz*, A Reynolds, AD Gaustein, GJ Peterson, JD Simmons, TR Hawn. University of Washington School of Medicine, Seattle, WA

Purpose of study Mechanisms that mediate natural resistance to Mycobacterium tuberculosis (Mtb) infection in highly exposed individuals are unknown. Using transcriptomic profiling of primary monocytes, we identified palmitic acid (PA)-induced genes that were enriched in donors with latent Mtb infection (LTBI) when compared to donors who resist infection despite high exposure. PA has known pleiotropic effects on macrophage responses including the activation of the NLRP3 inflammasome to increase IL-1β secretion. We hypothesized PA treatment of infected macrophages would impact the early mycobacterial response.

Methods used Intracellular BCG-cherry and Mtb-mlux replication within WT U937 macrophages was measured by fluorimetry and luminometry, respectively. Mtb-mlux growth was also measured in U937 cells with inactivating mutations in inflammasome genes (NLRP3 and NLRC4). IL-1β secretion from WT U937 cells was measured by ELISA.

Summary of results PA potentiated BCG and Mtb growth in WT U937 cells in a dose-dependent pattern. In contrast, oleic acid (OA) treatment did not modulate mycobacterial growth. Furthermore, Mtb whole cell lysate (TBWCL)-induced IL-1β secretion was increased in PA-conditioned WT U937 cells. However, Mtb growth potentiation by PA was similar in WT, NLRP3-deficient and NLRC4-deficient U937 cells.

Conclusions PA induced mycobacterial growth and IL-1β secretion in Mtb/BCG-infected and TBWCL-stimulated macrophages, respectively. However, the induction of IL-1β is consistent with known effects of PA on the inflammasome, alternative host pathways are presumed to be responsible for the PA-induced potentiation of mycobacterial growth since the NLRP3- and NLRC4-deficient cells had a similar phenotype as WT. Targeting these PA-dependent pathways with host-directed therapies could provide adjunctive treatments for tuberculosis.

329 VARICELLA ZOSTER VIRUS OPEN READING FRAME 9 DOMINATES SKIN TISSUE-RESIDENT MEMORY CD8 T CELL RESPONSE AFTER ZOSTER
MS Elder Waters*, VL Campbell, K Basu, S Mortazavi, R Tu, K Laing, C Johnston, DM Koelle. University of Washington, Seattle, WA

Purpose of study Varicella Zoster Virus (VZV) causes chickenpox during primary infection and establishes latency in dorsal root ganglia. Herpes zoster (HZ) occurs when VZV reactivates to cause a painful dermatomal rash. T cells specific for VZV are generated during VZV infection and are thought to be important for limiting HZ. Recently, a subset of T cells, tissue resident memory (TRM), have been described that do not recirculate after assuming residence in barrier tissues and are hypothesized to provide early surveillance against (re) infection. Placement of TRM is an aspirational goal for vaccines but little is known about their basic biology. In this study we measured if TRM persist in skin up to 1-year post-HZ and investigated which specific VZV proteins are most commonly recognized.

Methods used Subjects with HZ were recruited and skin biopsies taken 45 (n=18), 90 (n=12), and 360 days (n=5) after rash onset from healed HZ and control skin. Skin T cells were non-specifically expanded for functional assays. A panel of subject-specific artificial antigen presenting cells (aAPC) was created by transfecting Cos-7 cells with HLA cDNA corresponding to each subject’s HLA type, and co-transfected with each VZV open reading frame (ORF). After skin TRM were added, CD8 T cell activation was measured by IFN-g ELISA. Synthetic peptides internal to VZV confirmed reactivity to VZV ORFs.

Summary of results We readily detected skin VZV-specific CD8 TRM skin up to 1 year after zoster. The most commonly recognized VZV protein was ORF9, with reactivity detected in 75%, 42%, and 20% of samples on days 45, 90, and 360 respectively. Overall, skin TRM decreased over the first year, paralleling a decline in blood.

Conclusions VZV-specific skin CD8 TRM are detectable up to 1 year after HZ. For unknown reasons, the abundant tegument ORF9 protein is commonly targeted by TRM. Ongoing studies will determine if the same CD8 T cell clonotypes persist in skin over time and measure how T cell clonotypes are compartmentalized in skin versus blood. Ultimately, we hope that combined single cell mRNA and T cell receptor sequencing of skin-derived T cells will permit determination of the transcriptional signature of pathogen-specific skin TRM.

330 CHARACTERIZING CLINICAL MANIFESTATIONS OF CONGENITAL ZIKA SYNDROME IN NON-MICROCEPHALIC INFANTS
1JS Cranston*, 1SF Tiene, 1K Nielsen-Saines, 2M Moreira, 2V Vasconcelos, 2A Da Costa. 1University of Washington, Seattle, WA; 2Universidade Federal de Minas Gerais, Belo Horizonte, Brazil

Purpose of study The full spectrum of abnormalities due to antenatal exposure to Zika Virus (ZIKV) have not been delineated. We analyzed the various clinical manifestations in children with Congenital Zika Syndrome (CZS) from Rio de Janeiro, Brazil.

Methods used Newborns of women with nonspecific viral symptoms during pregnancy or referral to Instituto Fernandes Figueira (IFF) for suspected ZIKV were included. Vertical exposure to ZIKV was determined by PCR or IgM serology. For neurodevelopmental evaluation, Bayley-III Scales of Infant and Toddler Development was used.

Summary of results The study enrolled 296 newborns born during the ZIKV outbreak in Brazil. In utero exposure to ZIKV was confirmed in 219 cases. Among the non-microcephalic infants (N=161), 76.8% had failure to thrive (FTT) and 68.1% had a neurologic abnormality. Among non-microcephalic infants with Bayley III exam scores (N=112), head circumference was a significant indicator of abnormal language score (p=0.004) and developmentally delayed language score (p=0.011).
Conclusions ZIKV exposed infants without microcephaly suffer from congenital symptoms similar to those with microcephaly. Infant head circumference z-score at birth of non-microcephalic infants is significantly associated with neurocognitive development. Recognition of the various CZS phenotypes and spectrum of severity can help ensure early intervention.

**Abstract 330**

**Table 1** Outcomes according to microcephaly presence in ZIKV-Exposed infants

<table>
<thead>
<tr>
<th>Variable</th>
<th>Microcephalic infants (n=53)</th>
<th>Non-microcephalic infants (n=112)</th>
<th>All infants (n=165)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Average Head Circumference z-score at Birth</td>
<td>-0.3 (0.14)</td>
<td>0.05 (0.14)</td>
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**Abstract 331**

**Figure 2** Individual scores on Bayley III assessment according to significantly correlated early clinical findings

Abnormal neurodevelopment, auditory function, and neuromotor function prior to 3 months of age were found to be correlated to poor Bayley III outcomes (see table 1). -1 and -2 standard deviation are depicted by the red and gray dotted lines, respectively. The solid lines display values and the interquartile range.

**Abstract 332**

**The Use of Ceftaroline in the Treatment of Pediatric Infections with a Focus on Methicillin Resistant Staphylococcus Aureus (MRSA): A Literature Review**

**Purpose of study** Cefaroline, a novel cephalosporin was recently approved for treatment of community acquired bacterial pneumonia (CABP) and soft tissue infections (STIs). However, literature on the effectiveness of ceftaroline in treatment of serious infections due to MRSA is limited. The purpose of this study is to investigate the clinical effectiveness of ceftaroline in pediatric patients with serious infections, with a focus on MRSA infection.

**Methods used** An extensive literature search was conducted on PubMed and Google Scholar using keywords ‘ceftaroline’, ‘community-acquired pneumonia,’ ‘skin infection’ and ‘MRSA.’ Only studies or case reports of patients <18 years of age, diagnosed with serious infections which included evaluation of patients with MRSA infection were included.

**Summary of results** Of the 19 studies, 4 met our inclusion criteria (table 1 below). Ceftaroline was as effective as other

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**Abstracts**

**331 EARLY PREDICTORS OF NEURODEVELOPMENT IN INFANTS ANTENATALLY EXPOSED TO ZIKA**

1SF Tiene*, 1JS Cranston, 1K Nielsen-Saines, 2M Moreira, 2Z Vasconcelos, 3A Da Costa.
1David Geffen School of Medicine, Los Angeles, CA; 2Fundação Oswaldo Cruz, Rio de Janeiro, Brazil; 3Instituto Fernandes Figueira, Rio de Janeiro, Brazil

10.1136/jim-2019-WMRC.331

**Purpose of study** Identify early predictors of poor neurocognitive outcomes in non-microcephalic infants antenatally exposed to ZIKV, and evaluate their prognostic validity.

**Methods used** This study enrolled pregnant women with fever and rash during the ZIKV epidemic in Rio de Janeiro, Brazil from September 2015 through June 2016. Infants were also enrolled due to an abnormal prenatal ultrasound, confirmed antenatal ZIKV exposure, or clinical suspicion of CZS. We compared clinical findings identified prior to 3 months of age to neurocognitive outcomes, as measured by the Bayley III assessment after 6 months of age.

**Summary of results** Antenatal exposure to ZIKV was confirmed in 219 cases by positive maternal or infant PCR or IgM serology. 161 of these infants were non-microcephalic, 53 were microcephalic. Of the non-microcephalic infants, 112 received Bayley III exams after 6 months of age. Neuromotor abnormalities are correlated with abnormal motor scores (p=0.038, PPV=27.1%), and lower composite cognitive scores (p=0.048). Neurodevelopmental abnormalities are correlated with abnormal cognitive scores (p=0.044, PPV=31.3%). Auditory abnormalities are correlated with abnormal language scores (p=0.004, PPV=100%) and lower composite language scores (p=0.009).

**Conclusions** This study found that early neuromotor, neurodevelopmental, and auditory abnormalities are correlated to poor neurocognitive outcomes; however, the prognostic validity of these findings is poor and all infants that are antenatally exposed to ZIKV are at increased risk of developmental delay and should be monitored closely.

**332 THE USE OF CEFTAROLINE IN THE TREATMENT OF PEDIATRIC INFECTIONS WITH A FOCUS ON METHICILLIN RESISTANT STAPHYLOCOCCUS AREUS (MRSA): A LITERATURE REVIEW**

1,2RC Chow*, 2Z Bustami, 2S Afzal, 2R Durvasula, 2S Madireddy, 1T Gupta, 2B Afghani.
1University of California, Riverside School of Medicine, Riverside, CA; 2UC Irvine School of Medicine, Irvine, CA; 3CHOC Hospital of Orange County, Orange, CA

10.1136/jim-2019-WMRC.332

**Purpose of study** Cefaroline, a novel cephalosporin was recently approved for treatment of community acquired bacterial pneumonia (CABP) and soft tissue infections (STIs). However, literature on the effectiveness of ceftaroline in treatment of serious infections due to MRSA is limited. The purpose of this study is to investigate the clinical effectiveness of ceftaroline in pediatric patients with serious infections, with a focus on MRSA infection.

**Methods used** An extensive literature search was conducted on PubMed and Google Scholar using keywords 'ceftaroline,' 'community-acquired pneumonia,' 'skin infection' and 'MRSA.' Only studies or case reports of patients <18 years of age, diagnosed with serious infections which included evaluation of patients with MRSA infection were included.

**Summary of results** Of the 19 studies, 4 met our inclusion criteria (table 1 below). Ceftaroline was as effective as other
Abstracts

Abstract 332 Table 1 Use of ceftaroline in pediatric infections with focus on treatment of MRSA

<table>
<thead>
<tr>
<th>First Author, Year Published</th>
<th>Type of Study</th>
<th># of patients in experimental group (ceftaroline), age range</th>
<th># of patients in control group (other antibiotic), age range</th>
<th>Diagnosis</th>
<th># of patients with MRSA</th>
<th>Variable measured (treatment outcome)</th>
<th>% with successful treatment in experimental group (ceftaroline)</th>
<th>% with successful treatment in control group (other antibiotic)</th>
<th>Side effects</th>
</tr>
</thead>
<tbody>
<tr>
<td>Blumer, JL 2016</td>
<td>Prospective Randomized Control</td>
<td>30 (2mo-17yrs)</td>
<td>10, ceftriaxone plus vancomycin (2mo-17yrs)</td>
<td>CABP</td>
<td>1</td>
<td>Clinical response (improvement)</td>
<td>52%</td>
<td>67%</td>
<td>Anemia, Pruritus, Vomiting</td>
</tr>
<tr>
<td>Korczowski, B 2016</td>
<td>Prospective Randomized Control</td>
<td>107 (2mo-17yrs)</td>
<td>52, vancomycin or cefazolin (2mo-17yrs)</td>
<td>Bacterial skin infection</td>
<td>18</td>
<td>Clinical outcome (cure)</td>
<td>89%</td>
<td>96%</td>
<td>Diarrhea, Rash, Vomiting</td>
</tr>
<tr>
<td>Hall, AM 2018</td>
<td>Case Report</td>
<td>1 (12 y/o female)</td>
<td>N/A</td>
<td>Fasciitis and Bacteremia</td>
<td>1</td>
<td>Blood culture and clinical outcome (cure)</td>
<td>100% (1 case)</td>
<td>N/A</td>
<td>Not Reported</td>
</tr>
<tr>
<td>Williams, AW 2015</td>
<td>Case Report</td>
<td>2 (8 y/o males)</td>
<td>N/A</td>
<td>1. CA-MRSA bacteremia, thigh abscess, osteomyelitis</td>
<td>2</td>
<td>Blood culture and clinical outcome (cure)</td>
<td>100% (2 cases)</td>
<td>N/A</td>
<td>Not Reported</td>
</tr>
</tbody>
</table>

Conclusions Our review suggests that ceftaroline is effective and well tolerated in pediatric patients and may be considered as an adjunctive therapy in patients with MRSA infection. Larger controlled trials are needed to study the effectiveness of ceftaroline in MRSA infections.

CINEMATICAL PORTRAYAL OF INFECTIOUS DISEASES

1W Vu, 2B Holbrook, 3W Dehority. 1University of New Mexico, Albuquerque, NM; 2Scientific Technologies Corporation, Albuquerque, NM

10.1136/jim-2019-WMRC.333

Purpose of study Uptake of harmful activities (e.g. use of cigarettes and alcohol by children) is associated with viewing movies with these themes. To our knowledge, no studies have characterized the cinematic portrayal of infectious diseases. We hypothesized that over the last 4 decades, these films would feature an increasingly negative portrayal of the scientific/medical community.

Methods used 1567 films were identified using 149 infection-related search terms on the Independent Movie Database (IMDB), a validated tool for cinematic research. Synopses were reviewed on IMDB, Wikipedia, or American Film Institute (AFI) websites. Movies without synopses, without infectious diseases as a major theme, or without a domestic theatrical release were excluded. Year of release, box office data, media references, Academy Awards, genre, pathogen, themes, and portrayal of the scientific/medical community were assessed.

Summary of results 440 films (21.8% foreign-productions) released domestically from 1910 to 2018 were eligible. Drama (43.5%) and horror (37.4%) were the most common genres. Major themes included apocalyptic (27.3%), pandemics (20.1%), government conspiracies (19.8%), and bioterrorism (14.0%). The most common pathogens were viruses (53.4% of films), with human immunodeficiency virus most frequent (61 films) and fungi the least (7 films). Fictional viruses were increasingly depicted in modern movies in a linear regression model (p=0.0078), while bacteria (real/fictional) were more commonly featured in older movies (p=0.0012/p=0.0001). Films averaged 31.9 media references (median 4, range 0-655). Box office mean was $27,400,000 (median $2,856,712; range $0-$409,000,000). There were no significant relationships between media references and box office totals with ‘negative’ or ‘positive’ themes. 13 films won academy awards. Of films portraying the scientific/medical community, 69.0% depicted negative/mixed portrayals (e.g. unethical medical research, 15.5%). Negative themes were not more common over time when analyzed by decade in a logistic regression model (p=0.074).

Conclusions The scientific/medical community has been negatively portrayed in films depicting infectious diseases. Research on the ability of cinema to create mistrust towards the medical/scientific community is warranted.
Morphogenesis and malformations
Concurrent session
10:15 AM
Friday, January 24, 2020

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VARIANTS IN QRICH1- A NEW NEURODEVELOPMENTAL SYNDROME WITH VARIABLE HYPERMOBILITY, LEG PAIN, CONGENITAL HEART DISEASE AND MILD DYSMORPHIC FEATURES- REPORT OF A NEW PATIENT

1CJ Curry*, 2L Higa. 1UCSF Fresno, Fresno, CA; 2Community Regional Medical Center, Fresno, CA

10.1136/jim-2019-WMRC.334

Case Report Loss of function variants in QRICH1 have recently been reported in 5 patients (Ververi et al 2018 and Lui et al 2019). QRICH1 encodes the glutamine-rich protein 1, likely to be involved in apoptosis and inflammation. These 5 patients had mild developmental issues, 2 had leg pain with elevated CK’s, 2 had short stature with a mild chondrodysplasia and 1 had congenital heart disease. A relatively consistent craniofacial appearance was reported with thin upper lip and prominent nasal tip.

Our patient presented at 15 months with a large secundum ASD, mild developmental delays, gastroesophageal reflux and subtle dysmorphic features. A oligoarray was normal. She returned at age 11 with a history of hypermobility, migraine of the tongue and widespread asymmetric vascular cutaneous patches were noted. MRI showed a fusiform aneurysm of the internal carotid artery compressing the left optic nerve. A mosaic somatic overgrowth syndrome was suspected and NGS-based somatic tumor testing of 479 genes was performed on biopsied lesions. Previous case reports were reviewed.

Summary of results The missense variant in PDGFRB p.Y562C was detected in all somatic samples with variant frequencies (14%, 17%) but absent in blood. This variant was reported in only one case with the same phenotype. Interrogation of an inhouse database of 2.399 specimens revealed four other tumors with somatic variants in PDGFRB, including in myofibromas. We found that features of the mosaic condition can overlap with germline PDGFRB variants, sharing multiple features of Kosaki syndrome and Penttinen syndrome; also, aneurysms have been reported in a child with infantile myofibromatosis.

Conclusions Primarily used to identify cancer treatment, NGS-based somatic tumor testing may implicate this novel overgrowth condition by detecting low-level somatic mosaicism that would have been missed by traditional methods. The PDGFRB-related conditions represent a spectrum of manifestations. PDGFRB variants are also the first known genetic cause of these fusiform aneurysms, suggesting roles of aneurysm surveillance and therapies targeted to the pathway such as kinase inhibitors.

335
PDGFRB-RELATED OVERGROWTH SYNDROME WITH ANEURYSM: A SECOND EXAMPLE


10.1136/jim-2019-WMRC.335

Purpose of study Activating germline variants in PDGFRB have been associated with Kosaki overgrowth syndrome, infantile myofibromatosis, and premature aging (Penttinen syndrome).

Whether these conditions are distinct or overlapping entities is not clear. We report a patient with PDGFRB-related condition identified by next-generation sequencing (NGS)-based genetic testing.

Methods used A 26-year-old female with extensive unrevealing investigations for congenital differences in skin, soft tissues, and blood vessels presented with progressive left eye blindness. On exam, she looked older for age, with sparse hair and coarse facial features. The left leg and right arm were larger and longer than their counterparts. Nodular lesions of the tongue and widespread asymmetric vascular cutaneous patches were noted. MRA showed a fusiform aneurysm of the internal carotid artery compressing the left optic nerve. A mosaic somatic overgrowth syndrome was suspected and NGS-based somatic tumor testing of 479 genes was performed on biopsied lesions. Previous case reports were reviewed.

Summary of results The missense variant in PDGFRB p.Y562C was detected in all somatic samples with variant frequencies (14%, 17%) but absent in blood. This variant was reported in only one case with the same phenotype. Interrogation of an inhouse database of 2.399 specimens revealed four other tumors with somatic variants in PDGFRB, including in myofibromas. We found that features of the mosaic condition can overlap with germline PDGFRB variants, sharing multiple features of Kosaki syndrome and Penttinen syndrome; also, aneurysms have been reported in a child with infantile myofibromatosis.

Conclusions Primarily used to identify cancer treatment, NGS-based somatic tumor testing may implicate this novel overgrowth condition by detecting low-level somatic mosaicism that would have been missed by traditional methods. The PDGFRB-related conditions represent a spectrum of manifestations. PDGFRB variants are also the first known genetic cause of these fusiform aneurysms, suggesting roles of aneurysm surveillance and therapies targeted to the pathway such as kinase inhibitors.

336
IDENTIFICATION OF NOVEL CANDIDATE RISK GENES FOR MYELOMENINGOCELE WITHIN THE GLUCOSE HOMEOSTASIS AND FOLATE AND ONE CARBON METABOLISM NETWORKS

1C Baker, 2P Hillman, 2L Hebert, 3Hixon, 4Ashley-Koch, 5H Northrup*, 6K Au. 1Stanford University, Stanford, CA; 2McGovern SOM at UHealth Houston, Houston, TX; 3School of Public Health at UHealth Houston, Houston, TX; 4Duke University Medical Center, Durham, NC

10.1136/jim-2019-WMRC.336

Purpose of study Neural tube defects (NTDs) occur secondary to incomplete closure of the neural tube. Studies have shown an etiology with both genetic and environmental factors, but the underpinning of NTDs are yet to be elucidated. Two associated metabolic pathways are folate one carbon metabolism (FOCM) and glucose homeostasis/oxidative stress (GHOS). The purpose of this study is to apply WES data from a cohort of patients born with myelomeningocele (MM) to the hypothesis that analysis of damaging mutational burden will give insight into genes related to increased NTD risk.

Methods used Case data was generated with ~500 exomes of MM patients, split evenly between the ethnic groups with the highest birth prevalence in the United States: Mexican Americans and Caucasian Americans. Control data was obtained through the public Genome Aggregation Database (gnomAD).
Abstracts

Using an in-house bioinformatic pipeline, variants were filtered for quality and functional annotation. Variants included had an allele frequency <1% and are predicted to be damaging (stop gain, stop loss, canonical splice site, or combined annotation dependent depletion (CADD) phred score 20). Gene lists related to FOCM and GHOS were generated using gene ontology.

Summary of results We identified ~4000 qualifying variants in 1056/1469 genes, with 63 genes meeting nominal significance (p<0.05) for mutational burden (44 in the FOCM network, 22 related to GHOS, and 3 overlap). No genes reached mutation burden significance after applying Bonferroni correction. Overall, 64% of our population had at least one variant in a nominally significant gene and 28% had more than one.

Conclusions We present the largest WES cohort of NTD patients to date and propose using a large public database as a control. Interestingly, there was no overlap of implicated genes between ethnicities in either pathway. Only 6 genes have previously been implicated in neural tube defects, but 17 of them are expressed during human neural tube closure. Thus, we present 58 new genes as consideration for potential NTD risk.

WNT16 REGULATES VERTEBRAL SIZE AND PATTERNING DURING ZEBRAFISH DEVELOPMENT

1AR Cronnath*, 2Y Hsu, 3RY Kwon, 1CI Watson. 1University of Washington, Seattle, WA; 2Harvard Medical School, Boston, MA

Purpose of study WNT16 has a critical role in chronic diseases of the spine, as evidenced by its link to genetic osteoporosis susceptibility. However, the function of WNT16 in axial skeletal development remains elusive. To better understand the spinal roles of wnt16, we tested the hypothesis that Wnt16 mediates vertebral size and patterning during skeletal development.

Methods used All studies were approved by the University of Washington IACUC. CRISPR-based gene editing was performed by co-injecting zebrafish embryos with two guide RNA/Cas9 complexes (IDT) targeting wnt16. Somatic mutants were generated in a double transgenic background containing fluorescent reporters for osteoblast progenitors (sp7:EGFP) and canonical Wnt/b-catenin signaling. Imaging was performed at 12 or 26 days post fertilization (dpf) for somatic mutants. Germline mutants were stained with 0.2% calcein to visualize mineralization and imaged at 15 dpf.

Summary of results We performed in vivo imaging of osteoblast progenitors in tandem with Wnt/b-catenin signaling at stages of skeletal development (figure 1). Somatic mutants with mosaic loss of wnt16 exhibited reduced vertebral size at 12dpf as well as sporadic vertebral fusions (figure 1, p<0.05). At 26dpf, we observed increased sp7-derived fluorescence in the caudal spine (figure 2, p<0.05). Germline wnt16 mutants exhibited severe reductions in vertebral size (figure 3, p<0.05).

Conclusions These data indicate Wnt16 regulates vertebral size and patterning in zebrafish. Because wnt16 mutants exhibited decreased body size, it is possible that smaller centra in mutants may be secondary to body elongation defects. Somatic mutants also exhibited altered segmentation, while germline mutants did not, suggesting that although wnt16 is not required for spine segmentation, its mosaic expression influences the patterning process. In all, the role of wnt16 in vertebral development has future implications on the study and treatment of low bone mineral density diseases.

CHEDDA SYNDROME: A CASE REPORT AND REVIEW

1RD Clark*, S Ramanathan. Loma Linda University School of Medicine, Loma Linda, CA

Case report We report a patient with CHEDDA syndrome, a newly-described neurodevelopmental and multiple congenital anomaly disorder caused by a de novo missense variant in ATN1 at 12p13.31: c.3160C>T (p.His1054 Tyr). CHEDDA syndrome is an acronym for congenital hypotonia, epilepsy, developmental delay and digital anomalies.

A term male with BW 3515g was born after a pregnancy complicated by nausea, hyperemesis and premature rupture of membranes at 7 mo. He was admitted at 10 days for lethargy and inadequate intake. He had nevus flammeus, thin lips, long philtrum, pointy chin, fetal pads, mild finger contractures of both hands, overlapping toes and facial telangiectasias. He had mixed hyper/hypotonia, choreoathetoid movements and opisthotonic posturing with crossed ankles. Echocardiogram was normal. Brain MRI/MRS showed a thin corpus callosum. At 10 mos, he had a submucous cleft palate, bilateral strabismus, GERD and sleep apnea.

At age 3, he had febrile seizures. An EEG showed diffuse background slowing but no epileptiform focus. He ate only...
pureed foods. He had no speech, crawled, and pulled to a stand. He did not walk but manipulated a wheelchair expertly. He is being evaluated for autism. A trio WES (Ambry, 10/17) was negative. A diagnostic research trio whole genome test (UCLA, Dr. Stanley Nelson, PI, 7/19) identified the ATN1 variant.

In March 2019, Palmer et al. defined CHEDDA syndrome in 8 patients, 2 mo-9 yrs, with recognizable facies and variable congenital anomalies. The 7 surviving patients had hypotonia, limited speech and were nonambulatory. Five had seizures. One had choreoathetoid movements.

CHEDDA syndrome is caused by heterozygous de novo missense variants in exon 7 in the histidine-rich HX repeat motif (1050-1065) of ATN1. Other genes with this motif (RERE, AUTS2) also have neurocognitive phenotypes. ATN1 encodes atrophin-1, a nuclear transcriptional regulator important in embryogenesis of the CNS and heart. A triplet repeat expansion in exon 5 in ATN1 causes dentatorubral-pallidoluysian atrophy, a progressive neurodegenerative condition not associated with congenital anomalies.

CHEDDA syndrome, a distinct and recognizable pattern of anomalies, defines a second phenotype associated with a different functional domain in ATN1, a gene known to cause a progressive neurodegenerative disorder.

**Abstracts**

**MONOZYGOTIC TWINS DISCORDANT FOR CRANIAL DYSINNERVATION DISORDER: EVIDENCE OF VASCULAR DISRUPTION**

1CG Tise*, 1CDeFilippo, 1MR Ruzhnikov, 1D Stevenson, 1Stanford University, Stanford, CA; 2UC Davis MIND Institute, Sacramento, CA

**Purpose of study** Congenital cranial dysinnervation disorders (CCDDs) are a collection of rare, non-progressive neurogenic syndromes caused by developmental abnormalities of cranial nerves (CNs) resulting in primary or secondary dysinnervation. Included in this collection is Moebius syndrome, a disorder presenting with non-progressive 6th (abducens) and 7th (facial) CN palsies resulting in facial weakness and inability to abduct the eyes; other CNs may also be involved. Affected individuals often present as infants due to parental concern for lack of facial expression, excessive drooling, and strabismus. The etiology of Moebius syndrome is unclear, as most cases are sporadic. Some hypothesize it is due to an ischemic vascular event during embryonic development; however, autosomal dominant inheritance is described in a few families, suggesting a genetic contribution.

**Methods used** Chart review, physical examination, and literature review.

**Summary of results** Here we describe a pair of ex-37 week monochorionic diamniotic twin girls; one with normal growth and development, and one presenting at the age of 19 months with micrognathia, reduced facial expression, and poor feeding. MRI performed on the affected twin revealed severely hypoplastic or absent 4th (troclear) CN bilaterally, left 6th CN smaller than right, and bilateral hypoplastic 7th and 9th (glossopharyngeal) CNs. Exam was notable for mild facial palsy with decreased expression of the lower face and drooling; no obvious limitation of extracocular eye movement was noted. History, physical exam, and imaging are consistent with a diagnosis of a CCDD, suggestive of Moebius syndrome.

**Conclusions** The contribution of genetic risk factors underlying the etiology of CCDDs such as Moebius syndrome (e.g. heritability) likely varies among affected individuals due to the degree of impact and penetrance of the specific genetic variant(s). This presentation of monozygous twins discordant for a CCDD provides evidence in favor of a vascular disruption hypothesis and against that of a genetic etiology, as genetic variation between monozygous twins is assumed to be low.

**Neonatology general IV**

**Concurrent session**

**10:15 AM**

Friday, January 24, 2020

**340 GROWTH TRENDS OF OVER 2500 PREMATURE INFANTS FROM BIRTH TO EIGHT YEARS**

1-2 Barnard*, 1S Saladero, 1A Wells, 1-2 J Ryu. 1Rady Children’s Hospital San Diego, San Diego, CA; 2University of California, San Diego, San Diego, CA

**Purpose of study** Although numerous studies characterize the growth of preterm infants postnatally from birth to term, there is less information regarding expected growth for premature infants beyond this period. Prior studies with smaller cohorts of premature infants suggest differences in this population’s growth compared to the World Health Organization (WHO) and Center for Disease Control (CDC) growth charts. However, to the best of our knowledge, there are no studies assessing the growth of preterm infants with this large of a cohort. We aimed to compare postnatal growth trajectories for premature infants from birth to eight-years-old to the CDC and WHO growth charts.

**Methods used** This retrospective cohort study includes 2778 premature infants, born at 32 weeks (w) or less, who received care within Rady Children’s Hospital’s Network over a ten-year period. Data consisted of weight (Wt), height (Ht), head circumference, weight-for-length and BMI at each Rady outpatient encounter. Age-adjusted z-scores were calculated using the CDC growth standards for Wt and Ht. The effects of sex, GA, BW, correcting for gestational age (CGA), ethnicity and race on growth patterns are being assessed.

**Summary of results** Preliminary results show statistically significant differences in the growth trends of preterm infants subdivided by BW or GA. Using a t-test, infants born <27 w are significantly more likely to weigh less and be shorter (p<0.001) than those born >27 w after CGA. Infants with BW <750 g are significantly more likely to weigh less and be shorter (p<0.001) than those with BW ≥ 750 g after CGA. Interestingly, infants born 27-32 w and <750 g (older and smaller) are significantly more likely to weigh less and be shorter than those born <27w and ≥ 750 g (p<0.05).

**Conclusions** As more extremely premature and very low BW infants are surviving, there is a need for further assessment of this subpopulation’s expected postnatal growth. Interestingly,
the extremely premature and/or low BW infants appear to not demonstrate catch-up growth on the WHO and CDC growth charts even by eight years of age. Therefore, there is a need for the development of new growth charts for these populations.

**Abstracts**

**341 OPTIMAL GROWTH IN PRETERM INFANTS**

1H Lee, 2H Cohen, 1,3S Joaquina*. 1Stanford University, Stanford, CA; 2University of California Berkeley, Berkeley, CA

10.1136/jim-2019-WMRC.341

**Purpose of study** There is a knowledge gap regarding sociodemographic and hospital-level factors that lead to optimal growth in preterm, very low birth weight (VLBW) infants. Suboptimal early growth of an infant can impact long-term growth and early onset of adult metabolic and cardiovascular diseases. The objective of this study was to determine hospital- and patient-level factors that contribute to suboptimal postnatal growth among preterm and VLBW infants in California.

**Methods used** This is a population-based study including 21,332 infants with a birth weight <1500g or gestational age of 23-32 weeks registered in the California Perinatal Quality Care Collaborative in 2008-2016. Z-scores of birth weight and discharge weight were calculated by 2013 Fenton-growth-chart-calculator. Infants below the 10th percentile of Z-score difference of birth weight and discharge weight (ΔWeightZ) were flagged and analyzed for trends. A multivariate generalized linear model with the birth hospital as a random effect was used to analyze risk factors for <10th percentile ΔWeightZ.

**Summary of results** In this cohort, the average z-score at birth weight was -0.039 (SD: 0.845), average z-score at discharge weight was -1.04 (SD: 0.92), and average ΔWeightZ was -0.98 (SD: 0.75). The incidence of <10th percentile ΔWeightZ was highest in Hispanic infants (72%). In the multivariate analysis, we identified several significant risk factors and protective factors for <10th percentile ΔWeightZ. Hispanic infants were more likely to be <10th percentile of ΔWeightZ (OR: 1.32; 95% [CI]: 1.1, 1.6). Infants born in regional hospitals were less likely to be <10th percentile of ΔWeightZ (OR: 0.62; 95% [CI]: 0.41, 0.94). Infants born at an older gestational age were less likely to be <10th percentile ΔWeightZ (OR for a 1-week increase: 0.64; 95% [CI]: 0.62, 0.67). Infants with a higher Apgar Score at 5-minutes were less likely to be <10th percentile ΔWeightZ (OR for a 1-unit increase: 0.94; 95% [CI]: 0.92, 0.97).

**Conclusions** A risk factor for <10th percentile ΔWeightZ is Hispanic ethnicity. Protective factors for <10th percentile ΔWeightZ include birth at a regional hospital, birth at later gestational age, higher Apgar score at 5-minutes. Overall, attention to growth among different hospital characteristics and nutrition methods can facilitate targeted interventions in preterm and VLBW infants.

**Abstract 342 Table 1 Means±SD Fatty Acid%**

<table>
<thead>
<tr>
<th></th>
<th>Baseline</th>
<th>Week 2</th>
<th>Week 3</th>
<th>Week 4</th>
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</thead>
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<tr>
<td></td>
<td>SO</td>
<td>CO</td>
<td>SO</td>
<td>CO</td>
</tr>
<tr>
<td>DHA</td>
<td>4.3±1.3</td>
<td>5.2±1.0</td>
<td>3.3±1.1</td>
<td>4.8±1.1*</td>
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<tr>
<td>EPA</td>
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<td>0.7±0.5</td>
<td>0.4±0.2</td>
<td>1.5±0.5*</td>
</tr>
<tr>
<td>ARA</td>
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<td>21.8±1.1</td>
<td>17.2±2.0</td>
<td>17.7±1.9</td>
</tr>
<tr>
<td>Linoleic</td>
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<td>6.5±2.6</td>
<td>13.9±3.5</td>
<td>10.0±3.0*</td>
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</tbody>
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**342 THE EFFECT OF AN IV COMPOSITE LIPID ON POLYUNSATURATED FATTY ACIDS IN NEONATES**

E Kim*, S Goudarhan, N Li, KL Calkins. University of California, Los Angeles, Los Angeles, CA

10.1136/jim-2019-WMRC.342

**Purpose of study** Preterm infants and infants with gastrointestinal disorders use IV lipids as a source of fatty acids. Pure soybean oil (SO) emulsions are associated with low arachidonic (ARA) and docosahexaenoic acid (DHA). This study’s purpose was to compare fatty acid profiles in neonates who received SO or a composite (CO) emulsion containing fish oil.

**Methods used** This was a prospective study that included neonates <14 days of age with a birth weight <2 kg or gastrointestinal disorder and parenteral nutrition (PN) >14 days. Gas chromatography/mass spectrometry was used to measure fatty acid% in red blood cell (RBC) membranes. Linear mixed effects models and t-tests were used to compare fatty acids.

**Summary of results** In CO (n=27) and SO (n=69), the mean (±SD) gestational age was 31±6 and 31±5 weeks, respectively (p=0.7). PN duration was similar in SO and CO (31±27 vs.27±26 days, p=0.5). There was no significant difference in lipid dose between groups (p=0.2). At specific time points, DHA and eicosapentaenoic acid (EPA) were greater in CO vs. SO, while linoleic acid was lower in CO vs. SO. When CO was compared to SO, ARA was greater at week 4 only (Table). DHA decreased over time (-0.3%/week and -0.2%/week in SO and CO, respectively, p<0.01 for both) with no difference in the rate of change between groups (p=0.1). While EPA significantly increased over time in SO and CO (0.04%/week and 0.2%/week, respectively, p<0.01 for both), there was a difference between groups (p<0.01). ARA decreased over time in both groups (-1.1%/week and -1.0%/week in SO and CO, respectively, p<0.001 for both) with no difference in the rate of change between groups (p=0.7).

**Conclusions** When compared to SO, CO was associated with a greater DHA and EPA RBC%. While changes in ARA over time were similar for the two groups, CO had an increased ARA% at week 4. It remains to be seen if CO improves neonatal outcomes.
### Abstract 343

**BIOCHEMICAL DIFFERENCES IN PRETERM INFANTS LESS THAN 1250 GRAMS AT BIRTH FED HUMAN MILK WITH HUMAN MILK BASED FORTIFIER VERSUS BOVINE BASED FORTIFIED HUMAN MILK**

1M Chang, 1L Barton, 1T Lin, 1R Ramanathan, 1R Cayabyab*. 1LAC+USC Medical Center, Keck School of Medicine, University of Southern California, Los Angeles, CA; 2Miller Children’s and Women’s Hospital, Long Beach, CA

10.1136/jim-2019-WMRC.343

**Purpose of study** Comparison of laboratory values in preterm infants less than 1250 grams at birth fed with human milk with human milk-based fortifier (EHM) or bovine milk-based fortifier (HMF).

**Methods used** Retrospective data collection in preterm infants less than 1250g at birth admitted to NICU from January 2016 to May 2019 who were fed with EHM (26 cal/oz, protein: 0.025g/ml, potassium: 1.12mg/ml) or human milk fortified with HMF (24 cal/oz, protein: 0.025g/ml, potassium: 1.14mg/ml). Demographics and laboratory values were collected after full enteral feeding was tolerated for 3 consecutive days and off parenteral nutrition. Data was compared using chi square test or Wilcoxon rank sum test as appropriate.

**Summary of results** There were 53 preterm infants included in the study. Preterm infants fed EHM were of lower gestational age, however time to full enteral feeding was not significantly different between the type of fortification of human milk. (Table 1) Serum albumin and blood urea nitrogen (BUN) were significantly higher in preterm infants fed EHM. (Table 2).

**Conclusions** Preterm infants fed EHM resulted in improved nutritional indices after achieving full enteral feeding. This could potentially improve growth outcomes of ELBW infants at discharge.

### Abstract 344

**INTRAUTERINE GROWTH RESTRICTION AND LIVER DYSFUNCTION IN SMALL FOR GESTATIONAL AGE INFANTS**

MG Dominguez Garcia*, SR Sardeasi, M Durand, R Ramanathan. LAC+USC Medical Center, Los Angeles, CA

10.1136/jim-2019-WMRC.344

**Purpose of study** Evaluate the association between intrauterine growth restriction (IUGR) in small for gestational age (SGA) newborns and liver dysfunction in the first 48 hours of life.

**Methods used** Retrospective study of SGA newborns admitted to LAC+USC Medical Center Neonatal Intensive Care Unit between January 2010 and October 2018. Data were collected from our electronic medical records. Blood laboratory tests were obtained within the first 24 to 48 hours of life including: Aspartate aminotransferase (AST), alanine transaminase (ALT) protein, albumin, glucose, and bilirubin.

**Summary of results** From 207 SGA infants (birthweight <10th percentile), 39 patients were excluded from the study because of congenital infection, hypoxic-ischemic encephalopathy, <24 weeks of gestation, or major congenital anomalies. A total of 168 patients were evaluated (female 51%, Hispanic 53%, <37 weeks 71%, birthweight 1367±694 g, gestational age 32.1±4.9 weeks, Mean +/- SD). Sixty three percent of patients had increased AST values (>40 U/L), 57% had albumin level <3.5 g/dl, 25% had hypoglycemia (<45 mg/dL), and 31% of patients developed cholestasis (direct bilirubin >1mg/dL).

**Conclusions** Our preliminary findings indicate that 63% of SGA infants show signs of liver dysfunction within the first 24-48 hours of life and 31% developed cholestasis. We speculate that this could be related to undernutrition in the intrauterine environment.

### Abstract 345

**TREATMENT WITH LANSOPORAZOLE IMPROVES PREGNANCY OUTCOME IN MICE**

I’Iwatari, 1BL Le, 1CM Kracony, 1D Vijayakumar, 1R Kong, 1M Sinota, 1RJ Wong, 1DK Stevenson. 1Stanford University School of Medicine, Stanford, CA; 2University of California, San Francisco, CA

10.1136/jim-2019-WMRC.345

**Purpose of study** Using a computational drug repositioning approach that leverages data to uncover novel therapeutic uses for drugs already developed and evaluated for safety, we identified 83 that may be effective for preventing spontaneous preterm birth (sPTB). Of these, lansoprazole, a proton pump inhibitor (PPI) currently used for treating gastric ulcers, was identified with a strong profile. Thus, we evaluated the effect of lansoprazole on pregnancy outcome using a mouse inflammation model of fetal wastage.

**Methods used** Pregnant FVB mice were treated on E6.5–E8.5 with saline (IP), progesterone (P4, 80 mg/kg IM), mineral oil (P4 vehicle, IM), lansoprazole (PPI, 24 mg/kg IP), or 5% DMSO (PPI vehicle, IP).P4 served as a positive control as it is currently indicated for preventing sPTB. At E7.5, 2h after treatment, mice were given lipopolysaccharide (LPS, 100 μg/kg
### Abstracts

#### OUTCOMES OF LIGATION VERSUS CATHETER OCCLUSION IN VERY LOW BIRTHWEIGHT INFANTS

**T Rauser*, A Marshall, P Agrawal, P Jung, K Rawson, BM Gordon, D Deming. Loma Linda University School of Medicine, Loma Linda, CA**

**Purpose of study** Patent ductus arteriosus (PDA) remains a serious and controversial problem in preterm infants. We have previously reported that surgical ligation for closure of moderate-to-large PDA in very low birthweight infants is associated with increased prevalence of bronchopulmonary dysplasia (BPD). Another known complication of ligation is postoperative hemodynamic instability or post-ligation syndrome (PLS). Recently, an alternative method of PDA closure via transcatheter occlusion has been introduced. We describe our outcomes from this method.

**Methods used** We prospectively collected data of infants admitted to our NICU who were treated by ligation or occlusion after failing pharmacologic closure. Infants were <29 weeks gestational age, admitted prior to 30 days of age, and survived >7 days. We determined the prevalence of BPD, PLS, necrotizing enterocolitis (NEC), retinopathy of prematurity (ROP), intraventricular hemorrhage (IVH), and periventricular leukomalacia (PVL). We compared gestational age (GA), age at treatment, age at discharge, size of PDA, and left atrial to aortic diameter (La:Ao) ratio between the two groups. Chi-square and t-Tests were used.

**Summary of results** Our cohort contained 34 infants (21 – ligation; 13 – occlusion). The occlusion group had a lower prevalence of PLS and a trend for older age at treatment than the ligation group. There were no differences in GA, age at discharge, size of PDA, and La:Ao ratio between the two groups. There were no differences in rates or severity of BPD, NEC, ROP, IVH, or PVL between groups.

**Conclusions** These findings suggest transcatheter occlusion may provide a lower risk of hemodynamic instability compared to surgical ligation. The rates of other outcomes may be related to gestational age rather than method of closure.

#### OUTCOMES OF LIGATION VERSUS CATHETER OCCLUSION IN VERY LOW BIRTHWEIGHT INFANTS

**Concurrent session**

10:15 AM

Friday, January 24, 2020

#### ASSESSING SAFETY AND EFFICACY OF C1 ESTERASE INHIBITOR FOR PREVENTION OF DELAYED GRAFT FUNCTION IN DECEASED DONOR KIDNEY TRANSPLANT RECIPIENTS

**E Huang, A Vo, N Anmmanman, J Choi, K Lim, S Sethi, J Kim, S Kumar, R Najjar, SC Jordan*. Cedars-Shaif Medical Center, West Hollywood, CA**

**Purpose of study** Delayed graft function (DGF) is related to ischemia-reperfusion injury (IRI) and may be complement-dependent. Complement activation through the manose binding lecting pathway is a suspected mediator of IRI/DGF. C1 esterase inhibitor (C1INH) specifically inhibits the MBL pathway, thus may have utility in preventing IRI/DGF. We previously reported a randomized placebo-controlled trial that treatment with C1INH was associated with a shorter duration of delayed graft function, less dialysis and higher estimated glomerular filtration rate (eGFR) at one year. Here, we report longer-term outcomes from this trial.

**Methods used** This is a post-hoc analysis of a phase I/II randomized-controlled trial enrolling seventy recipients of deceased donor kidney transplants at risk for delayed graft failure. The full trial was published previously. We used the gift model of delayed graft function at 3 and 6 months post-transplant.

**Comparison between groups** Occlusion 25.3 ± 1.1 38 ± 17 118 ± 59 2.6 ± 0.8 1.5 ± 0.2

**Conclusions** Our cohort contained 34 infants (21 – ligation; 13 – occlusion). The occlusion group had a lower prevalence of PLS and a trend for older age at treatment than the ligation group. There were no differences in GA, age at discharge, size of PDA, and La:Ao ratio between the two groups. There were no differences in rates or severity of BPD, NEC, ROP, IVH, or PVL between groups.

**Conclusions** These findings suggest transcatheter occlusion may provide a lower risk of hemodynamic instability compared to surgical ligation. The rates of other outcomes may be related to gestational age rather than method of closure.
function (NCT02134314). Subjects were randomized to receive C1INH 50 U/kg (n=35) or placebo (n=35) intraoperatively and at 24 hours. The cumulative incidence functions method was used to compare graft failure and death over 3.5 years. eGFR slopes were compared using a linear mixed effects model.

**Summary of results** Seven graft failures developed in the placebo group compared to one among C1INH-treated recipients; the cumulative incidence of graft failure was lower over 3.5 years among C1INH-treated recipients compared to placebo (p=0.03). Although no difference in eGFR slopes was observed between groups (p for group-time interaction=0.12), eGFR declined in placebos-treated recipients (-4 ml/min/1.73 m²/year; 95% CI: -8 to -0.1) but was stable in C1 esterase inhibitor-treated patients (eGFR slope: 0.5 ml/min/1.73 m²/year; 95% CI: -4 to 5). At 3.5 years, eGFR was 56 ml/min/1.73 m² (95% CI: 42-70) in the C1 esterase inhibitor group vs. 35 ml/min/1.73 m² (95% CI: 21-48) in the placebo group, with an estimated mean eGFR difference of 21 ml/min/1.73 m² (95% CI: 2 to 41 ml/min/1.73 m²).

**Conclusions** Treatment of patients at risk for IR/DFG with C1INH at transplant was associated with improved eGFR and lower incidence of graft failure at 3.5 years. Thus, C1INH treatment at time of kidney transplant has the potential to improve the utility and function of marginal kidneys that are currently discarded.

**ADVANCED CHRONIC KIDNEY DISEASE IS ASSOCIATED WITH HIGHER AND NOT LOWER INSULIN USE**

1D Grube*, 2S Beddhu, 3R Boucher. 1University of Washington, Seattle, WA; 2University of Utah, Salt Lake City, UT

10.1136/jim-2019-WMRC.348

Purpose of study As kidneys metabolize insulin, it is commonly held that insulin use is lower with more advanced CKD. On the other hand, more advanced CKD might result in progressive loss of beta cell function, increase in peripheral insulin resistance and contraindications to other antidiabetic medications, all of which might increase the need for insulin. Better understanding of the relationship of CKD and insulin use will help to improve treatment in patients with type 2 diabetes and CKD.

Methods used We performed an observational study of 944,891 veterans with an ICD-9 code of type 2 diabetes mellitus and a serum creatinine lab drawn between 1/1/2008 and 12/31/2010. The incidence of baseline and subsequent insulin use was determined by the first time that the patient had a prescription for insulin after the index date in the VA database and was identified by pharmacy data and tracked until 12/31/2013. We measured urine concentrations of the kidney tubular injury biomarker kidney injury molecule-1 (KIM-1) and neutrophil gelatinase-associated lipocalin (NGAL) during the final week (week 5) of both low sodium (LS; target of 50 mmol/d) and normal sodium (NS; target of 150 mmol/d) intake. We previously demonstrated in that 5-weeks of dietary sodium restriction (DSR) reduces large-elastic artery stiffness as well as blood pressure in healthy middle-aged/older adults with moderately elevated systolic blood pressure and free from chronic kidney disease (CKD) as compared to normal sodium intake. Given these changes, we hypothesized that DSR would also reduce urinary concentrations of renal tubular injury biomarkers in this cohort, which are associated with more rapid loss of kidney function and predict incident CKD in the general population.

Methods used This was a post-hoc analysis using stored 24-hr urine samples collected in 13 participants as part of a randomized, double-blind, cross-over clinical trial of DSR. We measured urine concentrations of the kidney tubular injury biomarkers kidney injury molecule-1 (KIM-1) and neutrophil gelatinase-associated lipocalin (NGAL) during the final week (week 5) of both low sodium (LS; target of 50 mmol/d) and normal sodium (NS; target of 150 mmol/d) intake. We hypothesized that DSR would also reduce urinary concentrations of renal tubular injury biomarkers in this cohort, which are associated with more rapid loss of kidney function and predict incident CKD in the general population.

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excretion was reduced from 149±7 to 66±8 mmol/d during
week 5 of each condition. Despite having preserved
kidney function, participants had a 31% reduction in
urinary NGAL concentrations with just 5 weeks of DSR (LS: 2.8±0.6; vs.
NS: 4.2±0.8 ng/mL; p<0.05). Results were similar when
normalized to urinary creatinine, which did not change between
conditions. KIM-1 concentrations, however, were below the
detectable limit in all but one sample.

Conclusions DSR reduces evidence of kidney tubular damage
in adults with moderately elevated SBP who are free from
prevalent kidney disease. Reducing dietary sodium intake
should be evaluated as a strategy to prevent incident CKD as
well as slow progression of prevalent CKD.

350 COMPENSATORY RENAL GROWTH PREDICTS OUTCOME IN CHILDREN WITH A SOLITARY FUNCTIONING KIDNEY

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10.1136/jim-2019-WMRC.350

Purpose of study Congenital anomalies of the kidney are frequent
causes of childhood chronic kidney disease (CKD), however
not all experience a decline in their glomerular filtration rate (GFR). The challenge is to identify those at risk of developing
CKD. We hypothesized that in children with a solitary functioning kidney (SFK), kidney size at diagnosis and the extent of compensatory renal growth predict long term outcome,
and could be incorporated into a risk stratified model of care.

Methods used Using a retrospective cohort design, we
included cases with SFK due to unilateral renal agenesis (RA)
or multicystic dysplastic kidneys (MCDK) between the years
2000-2017. We extracted clinical data from medical charts;
sounderized kidney lengths (kidney length/body height) from
ultrasounds at first postnatal evaluation in the first year of
life, and used the last documented estimated GFR as the
primary outcome, with <90 ml/min/1.73 m² as our definition
of CKD. Between group differences were analyzed by t-test,
and receiver operating curve (ROC) analysis was used to predict
CKD.

Summary of results We identified a total of 229 SFK cases (71
RA, 158 MCDK) with median age of 0.3 (IQR 0.6) years at
initial evaluation and median follow up of 4.3 (IQR 4.8)
years. At last follow up mean eGFR was 88.9 ± 28.4 ml/min/
1.73 m², with a total of 53 cases (23%) with CKD. In those
initially evaluated in the first year (n=31), mean follow-up
was 4.3 ± 3.8 years, mean standardized length was
0.102±0.001, and 10 cases (32%) developed CKD. When
stratified by outcome, those who developed CKD had significantly
smaller standardized kidney lengths compared to those
who did not develop CKD (0.093 ±0.005 vs 0.106 ±0.003,
p=0.02). By ROC analysis, a standardized kidney length
below 0.101 optimally predicted CKD (AUC 0.722, 95% CI
0.514–0.931).

Conclusions In this study of children born with a SFK, a low
standardized kidney length at initial assessment, reflecting
impaired compensatory renal growth, was associated with the
development of CKD. These results will help us stratify long
term risk of CKD, will be used in the development of a clinical
pathway, and will allow the delivery of more focused and
efficient long-term clinical care.

351 EFFECT OF SHAM SURGERY AND UNILATERAL NEPHRECTOMY ON AUTOPHAGIC FLUX IN KIDNEY
AND HEART

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10.1136/jim-2019-WMRC.351

Purpose of study Compensatory renal hypertrophy resulting
from loss of nephron mass promotes nephron damage and
CKD. Unilateral nephrectomy (UNX) is a model of compensa-
tory renal hypertrophy in the remaining kidney. The purpose of
the study was to determine the effect of UNX on
mTORC1/2 signaling, lysosomal biogenesis and autophagic
flux in the kidney.

Methods used A model was developed to study autophagic
flux in vivo in the kidney and heart by treating mice with vehicle or bafilomycin A1 (BAF) and measuring LC3-II and p62 by immunoblot analysis. An increase in LC3-II (marker of
autophagosomes) after BAF or a decrease in p62 (degraded by
autophagy) was used as a marker of autophagy.

Summary of results On immunoblot analysis, sham surgery
(SS) and UNX resulted in increased p66 (mTORC1) and
pAkt (mTORC2) in the remaining kidney. mTOR is known to
regulate transcription factor EB (TFEB) and lysosomal bio-
genesis. LAMP2 protein expression and Lamp2, TFEB, and
Atg6a/d2 transcripts were decreased in SS and UNX indicat-
ing decreased lysosomal biogenesis vs. normal. As mTOR
and lysosomal function are crucial to autophagy, autophagic
flux was measured. There was a significant increase in LC3-II
in normal kidneys and heart 2 hours after BAF treatment. After
both UNX and SS, the increase in LC3-II after BAF was
absent indicating suppressed autophagic flux vs.controls. On
both IF and EM, the increase in autophagosomes was absent
after both SS and UNX vs. normal. Rapamycin inhibited the
increase in mTORC1/2 but did not restore autophagic flux
nor correct the lysosomal abnormalities. Autophagic flux in
heart was suppressed after UNX, but not sham surgery. To
determine whether other surgical procedures affect autophag-
ographic flux, a 24 hr bilateral renal ischemia/reperfusion(I/R)
model was performed in mice. LC3-II did not increase with
BafA1 in kidney after I/R vs. normal mice and SS. Unexpect-
edly, LC3-II did not increase in heart with BAF after both
SS or I/R.

Conclusions Increased mTORC1 and 2 signaling, lysosomal
abnormalities and suppressed autophagic flux were found in
the kidney after both SS and UNX. Both SS and I/R sup-
pressed autophagic flux in the heart. Suppressed autophagic
flux caused by sham surgery may have important implications
for interpreting results of autophagy studies using animal mod-
els requiring surgery.

352 CHRONIC KIDNEY DISEASE (CKD) PREVALENCE AND GLOMERULAR FILTRATION RATE (GFR) TRENDS IN
CHILDREN WITH TYPE 1 DIABETES (T1D)

K Favel*, C Panagiotopoulos, C Mammen. UBC, Vancouver, BC, Canada

10.1136/jim-2019-WMRC.352

Purpose of study Identification of CKD in pediatric T1D patients is important given their increased risk of end stage
renal disease (ESRD) in adulthood. However, literature analy-
zing the burden of CKD and GFR trajectory in pediatric
Acute interstitial nephritis is a rare and unusual side effect of omalizumab

ACUTE INTERSTITIAL NEPHRITIS A RARE AND UNUSUAL SIDE EFFECT OF OMALIZUMAB

Purpose of study Drug-induced acute interstitial nephritis (DI-AIN) accounts for 70-75% of acute kidney injury cases. Although any drug can cause AIN, there have been few reports of AIN caused by immune-modulator drug therapy. Here, we share the first reported case of AIN following administration of omalizumab for the treatment of moderate to severe persistent asthma.

Methods used Retrospective case report

Summary of results A 71-year-old female with moderate to severe asthma, uncontrolled type II diabetes, hypertension, heart failure with preserved ejection fraction, and chronic kidney disease stage four presented to the emergency department (ED) for progressively worsening shortness of breath and productive cough. She was in her normal state of health until three weeks prior to presentation when she received her first omalizumab injection for refractory asthma. After the injection, she developed mild generalized weakness and fatigue that worsened after her second injection two weeks later. In the ED, she was treated for fluid overload secondary to CHF exacerbation and renal failure, and acute asthma exacerbation. Subsequently, she underwent emergent dialysis as she had elevated levels of potassium of 8.1 mEq/L, BUN of 139 mg/dl, and creatinine of 8.08 mg/dl, suggestive of worsening kidney function as her baseline creatinine was 1-2 mg/dl prior to this admission. After two dialysis sessions, her symptoms significantly improved. She was then started on high-dose methylprednisolone for possible AIN secondary to omalizumab versus eosinophilic granulomatosis with polyangiitis. Her antinuclear antibodies screening returned positive but negative for antineutrophil cytoplasmic antibodies (ANCA) for myeloperoxidase and proteinase 3. A renal biopsy demonstrated no evidence of lupus nephritis or ANCA-associated crescentic glomerulonephritis with focal eosinophils noted near the cortico-medullary junction, suggestive of a DI-AIN. The patient’s hospital course remained uncomplicated and she was discharged with oral steroids and outpatient hemodialysis awaiting renal recovery.

Conclusions Our case highlights the first report of AIN as a side-effect of omalizumab and the importance of continued surveillance of rare side effects from current medications on the market.
febrile with recurrent partial small bowel obstruction, worsening peritonitis. Paracentesis was performed which revealed an elevated nucleated cell count of 15,000 and multiple loculations in her abdomen. Despite initial treatment with antibiotics the patient had 1-2 febrile episodes every day during admission. Over the course of the admission, she required multiple therapeutic paracenteses. Cachexia worsened and nutritional status deteriorated with albumin 1.1. Patient was started on TPN for symptomatic relief. Given deteriorating medical status, patient and family decided patient be placed on comfort care, and passed away on day 15 of admission.

**Conclusions** Encapsulating peritoneal sclerosis is a late complication of peritoneal dialysis characterized by intraperitoneal inflammation and fibrosis resulting in encasement of bowel. Despite trial of tamoxifen and prednisone, patient continued to deteriorate. Education about this rare complication of PD is crucial and further research for treatment for EPS is needed.

### Neuroscience II

**Concurrent session**

**10:15 AM**

**Friday, January 24, 2020**

**355** PROGNOSTIC VALUE OF THE UCSD PERFORMANCE-BASED SKILLS ASSESSMENT FOR COGNITIVE DECLINE IN PARKINSON’S DISEASE

1I Trang*, 1AE Simpson, 2LD Medina, 1BM Kluger, 1SK Holden. 1University of Colorado Anschutz Medical Campus, Aurora, CO; 2University of Houston, Houston, TX

**Purpose of study** In Parkinson’s disease (PD), functional impairments in cognitively demanding activities of daily living (ADLs) (e.g. paying bills, planning activities) can precede cognitive changes detectable by neuropsychological testing by up to 10 years. However, current screening protocols for cognitive decline do not objectively assess ADLs. The UCSD Performance-Based Skills Assessment (UPSA) is a 30-minute test for assessing ADLs that we recently validated in PD. We now examine the UPSA’s potential to predict cognitive decline at an earlier time point than standard cognitive screening tests.

**Methods used** 47 non-demented PD participants (table 1) completed the UPSA, neuropsychological battery, motor exam, and mood and quality of life scales at baseline and one year. Z-scores from neuropsychological tests were averaged into composite cognitive scores. Cognitive classifications were assigned by expert consensus conference.

**Summary of results** 11/47 (23.4%) PD participants converted to a more impaired cognitive class during the one-year follow-up. Baseline UPSA scores were not significantly different (p=0.16) between converters (x=79.4, SD 9.4) and non-converters (x=83.6, SD 7.8) and did not predict conversion (OR 0.95, p=0.24). However, lower baseline UPSA score predicted more rapid decline in the composite cognitive score in converters compared to non-converters (F(1,9)=10.1, p=0.01).

**Abstract 355 Table 1** Baseline characteristics of PD cohort

<table>
<thead>
<tr>
<th>Metric</th>
<th>Mean ± SD, Range</th>
</tr>
</thead>
<tbody>
<tr>
<td>Age (y)</td>
<td>68.9 ± 7.4, 53-83</td>
</tr>
<tr>
<td>Sex (% female)</td>
<td>34%</td>
</tr>
<tr>
<td>Education (y)</td>
<td>16.9 ± 2.8, 10-21</td>
</tr>
<tr>
<td>Disease Duration (y)</td>
<td>6.0 ± 7.2, 0.5-42.0</td>
</tr>
<tr>
<td>LEDD (mg/day)</td>
<td>522.7 ± 428.1, 0-1724</td>
</tr>
<tr>
<td>Cognitive Classification (%)</td>
<td>PD-Normal Cognition: 72.3% PD-Mild Cognitive Impairment: 27.7%</td>
</tr>
</tbody>
</table>

**Conclusions** The UPSA may identify those at risk of more rapid cognitive decline in PD. Our cohort will continue to be followed over time to allow for more robust exploration of cognitive functional decline in PD.

**356** ANNUALIZED RATE OF CORTICAL CHANGE IN FIRST EPISODE PSYCHOSIS

1,2K Dhaliwal*, 1P Lizano, 2OLutz, 3MKeshavan. 1Western University of Health Sciences, Pomona, CA; 2University of Colorado Anschutz Medical Campus, Aurora, CO; 3Northwestern University, Chicago, IL

**Purpose of study** First episode psychosis (FEP) studies evaluate patients during a critical time in which diagnosis and treatment intervention can improve a patient’s prognosis from complete recovery to development of psychosis spectrum disorders, including schizophrenia (SZ) or affective psychosis (NSZ). However, there are few identified structural and neurobiological markers that can help with early detection. The role of increased peripheral inflammatory states on neuroimaging has not been thoroughly evaluated.

We examined baseline and annualized rates of change (ARCH) structural differences between healthy controls (HC) and FEP patients and its association with clinical or inflammatory markers.

**Methods used** Baseline neuroimaging (FreeSurfer v6.0) and clinical measures were obtained for FEP-SZ (n=37), FEP-NSZ (n=23) and HC (n=47). ARCH was calculated between baseline and maximum timepoints for volume, surface area and thickness. General linear models assessed group differences. Spearman correlations were performed between cortical measures to clinical and inflammatory variables at baseline and with ARCH scores. Data was controlled for age, sex and race. Volume was also controlled for total intracranial volume.

**Summary of results** Baseline left anterior cingulate cortex (ACC) volume (d= -0.52, p=0.02) and surface area (d=-0.4, p=0.04) was lower in the FEP-SZ compared to the HC group. Left ACC thickness was lower (d=-0.56, p=0.03) in FEP-NSZ compared to FEP-SZ. Greater baseline IL10 levels was associated with lower total ventricle volume change (r=-0.44, p=0.05) and greater total volume (r=0.48, p=0.03). Within the FEP group, ARCH for left ACC volume, gray thickness and total gray volume were associated with decreased repetitive motion soft neurological signs.

**Conclusions** ACC plays a role in cognition, emotion and has been implicated in SZ pathogenesis. Evaluation of ACC is a potential predictive marker of psychosis risk and potentially a
prognostic marker of clinical outcome. IL10 may be protective against gray matter volume loss and ventricular volume enlargement.

PHENOTYPIC HETEROGENEITY IN AMYOTROPHIC LATERAL SCLEROSIS: A POTENTIAL EXPLANATION FOR THERAPEUTIC FAILURE
K Denson*. Medical College of Georgia, Augusta, GA
10.1136/jim-2019-WMRC.357

Purpose of study Current diagnostic criteria for amyotrophic lateral sclerosis (ALS) results in a heterogeneous patient population for clinical trials. Variability in patient presentation has resulted in a large spectrum of clinical signs that may represent distinct disease entities under the heading of ALS. This may explain why progress made in elucidating the etiology and pathophysiology of ALS has not been translated into effective treatments. The objective of this study is to utilize a new diagnostic scheme to stratify disease presentation.

Methods used Retrospective chart reviews were conducted on patients with ALS at Loma Linda University's ALS clinic between April 2013 and July 2019. Clinical presentation was categorized on four axes, depicting the extent of:

1. Upper motor neuron signs
2. Lower motor neuron signs
3. Bulbar involvement
4. Behavioral/cognitive involvement

We used heat map analysis to identify trends in phenotypic progression to determine if patient presentation was maintained and to compare clinical courses between subgroups. A Bowker’s test of symmetry was conducted to ascertain if overall change in phenotype was significant over three timespans. Patients were omitted if they presented to the clinic over a timespan of less than 9 mos. and if enough data was not presented to determine the axes. The cognitive axis was not determined due to inconsistency of data.

Summary of results Seven distinct phenotypic subgroups were identified:

1. Upper motor neuron
2. Lower motor neuron
3. Bulbar
4. Upper and lower motor neuron
5. Lower motor neuron and bulbar
6. Upper motor neuron and bulbar
7. Generalized

Of 28 patients presenting with lower motor predominance, 21 (75%) remained lower motor predominant and 28 (100%) had lower motor neuron involvement one year after initial presentation. Patients with bulbar predominance became more generalized between 9 mos. and 12 mos. (100% vs 45.5% remaining bulbar predominant, respectively). There was no significant change in phenotype from first visit to 9 mos., first visit to 12 mos., or 9 mos. to 12 mos. using an adjusted significance level of 0.0167.

Conclusions Preliminary data suggests that there are phenotypic subgroups within the spectrum of ALS that remain consistent in the disease course. These data provide justification to further explore distinct pathophysiology and treatment responses in these groups.

PATIENT DEMOGRAPHICS AND DISEASES CHARACTERISTICS PREDICT LIKELIHOOD OF IMPROVEMENT ON PATIENT-REPORTED OUTCOME MEASURES IN MULTIPLE SCLEROSIS
JB Leary*, S Silfau, BL Vollmer, K Nair, TL Vollmer. University of Colorado Health Sciences Center, Aurora, CO
10.1136/jim-2019-WMRC.358

Purpose of study Patients with multiple sclerosis (MS) are increasingly treated with high-efficacy disease-modifying therapies (DMTs). Clinicians have observed functional improvement in a subset of these patients, but the factors influencing improvement are unknown. We examined the impact of demographics, disease characteristics, and brain volumetrics on the likelihood of clinical improvement in patients treated with high-efficacy DMTs, as assessed by patient-reported outcome (PRO) measures.

Methods used This retrospective chart review included adults with MS (>18 years) who had completed ≥2 PRO measures across 2 time points separated by ≥10 months and taking fingolimod, dimethyl fumarate, natalizumab, rituximab, or ocrelizumab at baseline. We examined the influence of patient demographics, disease characteristics and brain volumes on the likelihood of clinical improvement. PRO measures included 10 domains of the NeuroQOL Short Form battery. Patients were grouped as Improved vs. Failed to Improve by each NeuroQOL domain, using ≥ standard deviation as the benchmark for clinically significant improvement. Statistical analyses utilized Spearman correlations and logistic regression.

Summary of results 318 patients met inclusion criteria. The sample was 75% female, average age of 47.1 years, 82% relapsing-remitting MS (RRMS), and mean time between PRO 1 and PRO 2 of 2.4 years. Factors significantly predicting the likelihood of clinical improvement and the relevant Neuro-QOL domain included: non-white race and Fatigue (Odds Ratio=2.325, CI 5, 95=1.122, 4.817, p=0.023), younger age and Depression (odds ratio=0.976, CI 5, 95=0.952, 1.000, p=0.049), and secondary progressive type of MS and Sleep Disturbance (PPMS v. SPMS odds ratio=0.101, CI 5, 95=0.012, 0.842; RRMS v. SPMS odds ratio=0.346, CI 5, 95=0.166, 0.721, p=0.0069).

Conclusions Demographics and disease characteristics appear to be stronger predictors of clinical improvement than brain volumes. Most patients improved or maintained function across domains, suggesting that high-efficacy DMTs may prevent further debilitation and enhance the likelihood of functional improvement.

EXPLORING RELATIONSHIPS BETWEEN NEURAL ACTIVATION AND THE AUTOMATICITY OF POSTURAL CONTROL
1 2EI Bent*, 3SM Monfort. 1Montana State University, Bozeman, MT; 2University of Washington School of Medicine, Seattle, WA
10.1136/jim-2019-WMRC.359

Purpose of study Postural control studies are widely used to both assess fall risk and reduce falls. An important variable in these studies is Sample Entropy (SEn), which measures the regularity of changes of the Center of Pressure (CoP) under the feet. Increased SEn, which is commonly associated with
more automatic postural control, has been observed in the presence of cognitive distractors. However, the direct link between SEn and neural activation has not been studied. This study identified relationships between neural activation and SEn during multiple balance conditions.

**Methods used** 30 subjects stood on a balance plate and wore a prefrontal cortex-directed functional near-infrared spectrometry (fNIRS) cap while undergoing four bouts of 30-second stance during four balance tasks: (1) EOST - eyes open single task (standing still); (2) EODT - eyes open dual task (standing still while counting back by 7’s); (3) ECST - eyes closed single task; and (4) ECDT - eyes closed dual task. fNIRS data were characterized with autoregressive iterative least squares beta values, then analyzed with a mixed effects model. CoP was measured using a balance plate (1000 Hz), and SEn was calculated for increment resultant CoP data.

**Summary of results** Preliminary SEn data suggest significant increases in SEn for eyes open conditions when compared to eyes-closed conditions. Significant differences in SEn were observed between EOST and ECST (p=0.002) and between EOST and ECDT (p=0.011). fNIRS data analysis is currently underway and will be completed before WSMBF. These fNIRS beta values will then be correlated to changes in SEn relative to the EOST condition.

**Conclusions** The differences in SEn between EOST and ECST, as well as EOST and ECDT, support our expectations that eyes-closed conditions are associated with more attentional investment in balance and a subsequent decrease in SEn. We expect that increased prefrontal cortex activation during ECST will be positively correlated with decreased SEn during ECST. This finding will support the interpretation of SEn being associated with attentional effort and the automaticity of postural control. Furthermore, increased prefrontal cortical activation during dual task conditions may be due to a mix of attentional investment and cognitive distractors.

### 360 A RARE CASE OF UNILATERAL MYELITIS IN A 69 YEAR OLD FEMALE: A CASE REPORT

Y. Jiao, M. Arias*, R. Shah, V. Podduturu, L. Huang, T.aylor College of Medicine, Houston, TX; Burrell College of Osteopathic Medicine, Las Cruces, NM; Enloe Rehabilitation Hospital, Chico, CA; Michael E. DeBakey Veterans Affairs Medical Center, Houston, TX

Transverse myelitis (TM), a rare inflammatory disease affecting the spinal cord, causes bilateral weakness, sensory deficits, bowel bladder issues and autonomic dysfunction. TM is diagnosed on the basis of radiographic and clinical findings. It is crucial to consider a wide range of etiologies including: neuro-inflammatory processes such as Neuromyelitis Optica Spectrum Disorder and MOG Antibody Associated Disease, vascular process such as spinal cord infarct, infectious process such as HIV and syphilis, connective tissue disorder such as lupus or scleroderma, nutritional deficiencies such as B12 or copper, or paraneoplastic syndrome. Below we present a unique case of idiopathic unilateral myelitis with correlating findings on imaging.

**Case presentation** A 69 years old Caucasian female who presented with acute onset of right-sided paralysis. Patient reports symptoms had insidious onset over 2-3 hours. She also complained of urinary incontinence. No sensory deficits, no bowel incontinence, and no reported visual symptoms. Examination of her cranial nerves were unremarkable. Sensory intact bilaterally. Motor exam reveal strength intact in left upper and lower extremities and paralysis of right upper and lower extremities. T2 weighted MRI spine with an area of hyperintensity on the right side of the spinal cord spanning from levels C3-C5. MRI brain without acute intracranial events. CSF analysis with increased proteins, no oligoclonal bands present. NMO-IgG and MOG-IgG antibodies both negative. Patient also underwent extensive laboratory workup of West Nile, CSF, thyroid, vitamin B12, vitamin E, MMR, meningitis, and complement 3 and 4, all negative except for C4 which was moderately elevated. Patient was diagnosed with idiopathic transverse myelitis and started on IV Solumedrol. Patient underwent a comprehensive rehabilitation program with PT, OT and physiatry with remarkable recovery. Motor recovery was noted to be distal to proximal. Follow-up visit 3 months later patient was able to ambulate using right lower extremity posterior leaf spring AFO with cane.
increases in osteoclast and inflammatory genes (RANKL, IL-6) in the morphine treated group; not seen in the JWH015 group. 

**Conclusions** Non-psychotropic cannabinoid therapies (i.e., CB2 agonists) may be more effective angesics for bone fracture that lack unwanted side effects and enhance bone healing.

**Purpose of study** Microglia are the resident immune cells of the brain responsible for protecting the central nervous system (CNS) against various pathogenic factors. Besides being involved in the immune response, microglia are highly dynamic, playing an important role in synaptic plasticity through synaptic pruning via phagocytic mechanisms. Although microglial role in synaptic plasticity and cognitive function has been well characterized, the impact of learning experience on microglia regeneration after microglia injury is unclear. To understand how learning and memory affect microglia regeneration, we chemically induced microglial death in a PLX-treatment mouse model and tracked elimination and repopulation of microglia in distinct layers of the hippocampus, a brain region critical for learning and memory.

**Methods used** Survival and development of microglia depend on colony-stimulating factor 1 receptor (CSF1R). PLX5622 is a CSF1R inhibitor well-studied in microglia elimination experiments. In our experiments, two groups of mice, the naïve group and learning group, were fed a PLX5622 containing diet (1200mg/kg) throughout the experiment. All mouse brains were extracted after completion of the learning experiment, sliced with vibratome, and probed with Iba1 antibody. Microglial density in different hippocampal CA1 layers was counted manually in a double-blinded fashion from fluorescence microscopic images.

**Summary of results** The majority (>90%) of microglia in the Stratum Pyramidale (SP) and Stratum Radiatum (SR) layers of hippocampus were eliminated in control mice after over three weeks of PLX5622 treatment. Mice in the learning group had significantly higher numbers of microglia both in SP and SR layers when compared to control mice. 

**Conclusions** The results from our study showed the mice exposed to the learning paradigm have a higher density of microglia throughout the hippocampus CA1 subregion. This indicates that learning has a positive effect on microglia regeneration. Further mechanistic studies will help elucidate if the accelerated regeneration of microglia will have a positive effect on memory in hopes of translating this information to humans suffering from cognitive disorders.

**Purpose of study** In the United States, 1 in every 6 Americans self-report experiencing migraine headache over a 3-month period. A major diagnostic criteria for migraine headache is photophobia (an abnormal sensitivity to light). Up to 80% of migraine patients experience photophobia. Due to the high prevalence of photophobia in migraine patients, it is imperative to have an effective method of measuring the severity and impact of photophobia in order to improve patient care for patients with migraine. Our group developed a new questionnaire including questions on interictal light sensitivity and ADLs. This questionnaire shows promise; however, it is not yet validated as a disease-specific questionnaire for migraine headache and requires additional validation.

**Methods used** Patients were screened in Headache and Neuroophthalmology clinic at the University of Utah for migraine headache. All patients completed the Headache Impact Test (HIT-6) and the Migraine Disability Assessment Test (MIDAS) as well as the Utah Photophobia Symptom Impact Scale (UPSIS-12). Internal consistency, construct validity, and difference of means were assessed.

**Summary of results** 109 patients were included. (60 migraine, 49 non-migraine). Good reliability was observed in migraine patients. (Cronbach’s $\alpha=0.87$) UPSIS-12 scores moderately correlated with HIT-6 scores. ($r=0.37, p=0.004$) An unpaired two sample t-test was conducted to determine a difference in mean UPSIS-12 scores between migraine patients and non-migraine patients. There was a significant difference of means between migraine patients (M=33.72, 95% CI=[29.63, 37.80]) and non-migraine patients. (M=18.48, 95% CI=[14.46, 22.50]).

**Conclusions** The UPSIS-12 is a valid and reliable measure of light sensitivity in migraine headaches. Future studies are needed to identify thresholds for different sub-types of migraine (Chronic migraine, Episodic, etc.) and future disease specific validation is necessary for other headache groups (cluster headaches, Post traumatic headache).

**Surgery IV**

**Concurrent session**

**10:15 AM**

**Friday, January 24, 2020**

**Abstracts**

**364 MISSED CHEMOPROPHYLAXIS DOES MATTER: A MULTICENTER ANALYSIS OF VENOUS THROMBOEMBOLISM CHEMOPROPHYLAXIS IN TRAUMATIC BRAIN INJURY PATIENTS**

1 Zangara,* 1 Coleman, 2 Carmichael, 2,3 Saussa, 2,4 Moore, 5 Dunn, 5 Schroepel, 4 Campion, 3 Goodman, 3 Floen, 2 Ferrigno. 1 University of Colorado School of Medicine, Aurora, CO; 2 University of Colorado-Denver, Aurora, CO; 3 University of Colorado-Denver, School of Public Health, Aurora, CO; 4 Ernest E Moore Shock Trauma Center, Denver, CO; 5 UCHealth Memorial Hospital, Colorado Springs, CO; 6 University of Cincinnati Medical Center, Cincinnati, OH; 7 University of Arizona, Phoenix, AZ

**Purpose of study** Traumatic brain injury (TBI) is associated with morbidity/mortality in trauma patients. Concern for progression of intracranial hemorrhage leads to delayed/missed doses of venous thromboembolism (VTE) chemoprophylaxis
### Abstracts

(VTE-C) and to higher risk of thrombotic complications. Consensus about utilization of VTE-C in TBI patients (TBI-P) is lacking and we hypothesize late/missed VTE-C in TBI-P results in increased VTE.

**Methods used** This was a multicenter, two-year study of adults with head abbreviated injury scale ≥2, ≥2 head CTs, and LOS ≥72h. We compared early vs late (48h) VTE-C initiation and proportion with missed V those without missed doses.

**Summary of results** Of 1,803, 8% developed VTE. Median age was 55, 69% were male, and 95% had blunt injury. Median time to VTE-C initiation was 63h. VTE-C was not received in 29%; late in 47%; initiated within 48h in 25%; and initiated within 24h in 3%. Of 1,288 receiving VTE-C, 1,092 missed no doses. In patients with missed doses (196), there was longer time to initiation (57 v 63h, p=0.02) and they were more severely injured (ISS 27 v 22, p<0.001). They had more polytrauma (22% v 9%, p<0.001), higher rates of VTE (20% v 8%, p<0.001), longer hospital LOS (17 v 8 days), and longer ICU LOS (12 v 4d), (both p<0.0001). There was no significant reduction in VTE with early VTE-C for the group as a whole, however, among TBI-P who underwent delayed neurosurgical intervention, early VTE-C was protective against VTE (HR 0.81, 95% CI 0.73–0.91). After multivariable regression, missed VTE-C doses significantly predicted development of VTE (HR 1.03, p=0.01).

**Conclusions** VTE-C is often delayed >48 h or held in TBI-P, predicting higher VTE risk. Future studies are needed to examine the safety of early and perioperative VTE-C.

<table>
<thead>
<tr>
<th>Abstract</th>
<th>Title</th>
<th>Authors</th>
<th>Summary</th>
</tr>
</thead>
<tbody>
<tr>
<td>10.1136/jim-2019-WMRC.365</td>
<td>QUANTIFICATION OF BONE VISCOELASTICITY WITH LAG SCREW IN HINDFOOT ARTHRODESIS</td>
<td>BW Johnson*, A Rugg, M Requist, A Alvarez, M Son, L Latt. The University of Arizona, Tucson, AZ</td>
<td>Non-union and delayed union, common complications following hindfoot arthrodesis, may be caused by failure to obtain and maintain compression across bone-to-bone interfaces. Partially threaded cannulated ‘lag’ screws are commonly used for fixation in hindfoot arthrodesis; however, the ability of these screws to generate and maintain compression across hindfoot bones has not been quantified. The goal of this project was to quantify the stress relaxation response of hindfoot bones with initial and repeated lag screw compression.</td>
</tr>
<tr>
<td>10.1136/jim-2019-WMRC.366</td>
<td>PROVIDONE-IODINE DOES NOT AFFECT ACELLULAR DERMAL MATRIX INTEGRATION IN PATIENTS UNDERGOING TWO-STAGED, PREPECTORAL, BREAST RECONSTRUCTIVE SURGERY</td>
<td>E O’Rourke*, RM Gold, M Pfaffenberger, E Eldenburg, T Storm-Dickerson, A Gabriel. Elson S. Floyd College of Medicine, Spokane, WA; Loma Linda University Medical Center, Loma Linda, CA; Private Practice Allen Gabriel MD, Vancouver, WA; Compass Oncology, Vancouver, WA</td>
<td>Purpose of study Providone-iodine has been used as an antimicrobial agent to irrigate the breast pocket and rinse prosthesis prior to placement in breast reconstruction surgery. Use of providone-iodine with breast implants was banned from 2000-2017. During this time, acellular dermal matrix (ADM) was introduced to breast surgery. In nonclinical studies, providone-iodine was shown to impair collagen synthesis and kill fibroblasts. As cellular repopulation is critical for ADM integration, concerns were raised about the impact of providone-iodine on ADM integration. However, whether or not providone-iodine impacts ADM integration is unknown.</td>
</tr>
</tbody>
</table>

**Methods used** Twelve pairs of 25mm-diameter bone cylinders were cut with a keyhole saw from fresh thawed cadaveric feet. After cartilage removal and subchondral flattening with an oscillating saw, a calibrated Futek LTH 300 donut load cell and two metal washers were sandwiched between cylinders. Data was filtered using a 6th order Butterworth filter with a 0.035 normalized frequency cutoff. An 8.0mm partially threaded cannulated lag screw (Smith and Nephew, Memphis, TN) was placed from the posterior calcaneus to the anterior-superior talus or posterior-inferior talus to inferioribia to simulate arthrodesis. Compression between cylinders was recorded at 10Hz as screws were tightened by three quarter-turns, rested for 3 minutes, retightened one quarter-turn, and rested for 30 minutes.

**Summary of results** Compression with a lag screw demonstrates increased maximum compression and slowed decay upon screw retightening. Maximum compression after three quarter-turns and retightening averaged 284 N and 351 N, respectively (p-value: 0.0136). Time to decay to 80% of maximum compression after three quarter-turns and retightening averaged 40 (n=9; SD: 57) and 528 (n=9; SD: 590) seconds, respectively. Compression loss 30 minutes after retightening averaged 25.3% (SD: 8.8%).

**Conclusions** The increase in maximum compression and slowed decay upon retightening with a lag screw can be applied in surgical technique before wound closure to increase the likelihood of successful arthrodesis. Lag screw compression also resulted in a significantly lower percent compression loss and loss rate compared to direct compression. Limitations of this study include its small sample size and the use of cadaveric bone. Future experiments will include a larger sample size and other types of screws.
ADM integration was noted in 97% of breasts in each group. At implant exchange, integrated matrices appeared healthy, had no signs of foreign body reaction, and demonstrated punctate bleeding. Complications did not differ between groups, including rate of infection, seroma, and expander loss.

**Conclusions** Providone-iodine used for irrigation of the breast pocket and presoaking of prosthesis and ADM appears to have no adverse effects on clinical outcomes. It also did not impede matrix integration. Providone-iodine is safe for irrigation of the breast pocket and presoaking of the prosthesis and ADM in breast reconstruction.

### Abstracts

#### 367 BACK (DOOR) TO THE FUTURE: DORSAL LUMBOTOMY FOR PEDIATRIC UPPER POLE HEMI-NPHERECTOMY

1A Roshan, 1AE Machielli, 1University of British Columbia, Vancouver, BC, Canada; 2BC Children’s Hospital, Vancouver, BC, Canada

10.1136/jim-2019-WMRC.367

**Purpose of study** The dorsal lumbotomy approach to renal surgery has become a lost art. Upper pole heminephrectomy (UHN) is performed for two main indications: ectopic ureter-ocle and duplication anomalies with upper pole ectopy. Current popular techniques for conducting UHN include open flank, laparoscopic, and robotic. This study evaluates outcomes following dorsal lumbotomy (DL), an open approach used historically for pyeloplasty and pyelolithotomy, and in which no clinical trials or exclusive case-series have been conducted for UHN in children.

**Methods used** A retrospective review of 50 UHN performed in pediatric patients using the DL approach by a single surgeon at BC Children’s Hospital between 2000–19. Clinical variables and indicators included age, sex, weight, skin to skin time, total operative time, duration of hospital stay, post-operative complications, analgesic requirements, and 3-month post-operative ultrasound results.

**Summary of results** Mean age at surgery was 24.5 months. Mean length of follow-up was 22.5 months. Mean (range) for time between skin incision and closure was 90 (62–140) minutes, and the mean (range) total operating room time was 140 (70–180) minutes. There were neither intraoperative complications nor transpositions. The mean (range) post-operative opioid delivered was 0.74 (0.00–2.00) mg/kg/day. Mean (range) post-operative ibuprofen delivered was 4.73 (0.00–25.00) mg/kg/day. Median length of hospital stay was 2 days. No patients received postoperative prescriptions for narcotics at discharge. Two patients experienced minor wound complications. One patient had secondary atrophy of the lower pole. Secondary lower tract surgery, unrelated to surgical approach, was performed in six patients. Fifteen patients experienced a urinary tract infection at some point after surgery.

**Conclusions** DL is a historical approach for UHN that should not be forgotten. It is safe, feasible, and produces operative outcomes and times comparable or superior to that of conventional open flank incision, laparoscopic, and robotic techniques. These findings as well as cost considerations should be considered when promoting robotic approaches to UHN.

#### 368 INCIDENCE OF PNEUMOTHORAX AS A COMPLICATION OF BREAST SURGERY

1AG Kulinich*, 1JR Mark, 1J Osborn, 1,2LL Pu, 1University of California Davis School of Medicine, Sacramento, CA; 2University of California Davis, Sacramento, CA

10.1136/jim-2019-WMRC.368

**Purpose of study** Pneumothorax is a complication of breast surgery which was thought to be rare until Osborn and Stevenson surveyed members of the California Society of Plastic Surgeons on the number of complications of pneumothorax in breast surgery and variables involved in their cases. The response rate of the survey was 50% and the incidence of pneumothorax seemed to be more common than generally appreciated with 1 out 3 surgeons experiencing at least one pneumothorax in their career. To our knowledge, no study has examined the incidence of pneumothorax in breast augmentation surgery procedures since the Osborn and Stevenson survey published in PRS. (Plast. Reconstr. Surg. 116: 1122, 2005) To address this research gap, we are performing a systematic review of the literature to evaluate the rates of complication of pneumothorax related to breast augmentation surgery and researching all institutions that may have worthwhile statistics on the subject.

**Methods used** A thorough search of the literature was conducted using the terms ‘pneumothorax’ and ‘breast augmentation’ in the PubMed/MEDLINE databases. The search generated 23 articles. This number was brought down to 6 after initial screening by title. Inclusion criteria then removed those not written in English, those without access to the full text, those without extractable data on complications, those older than 10 years and duplicates, leaving 3 case reports to examine between 2008 and 2019.

**Summary of results** The incidence of pneumothorax in breast surgery appears, at this time, to be in the range of 0.1% to 0.53%. This is consistent with the findings from our previous survey.

**Conclusions** Pneumothorax is still a rare complication after breast augmentation. Our findings confirm the conclusions of our earlier article regarding the encouragement to include pneumothorax as a risk of breast augmentation in the informed consent.

#### 369 HOW GOOD IS THE MUSTARDE OTOPLASTY?

1M Baroščik*, 1AV Vanslyke, 12JS Arneja, 1University of British Columbia, Vancouver, BC, Canada; 2BC Children’s Hospital, Vancouver, BC, Canada

10.1136/jim-2019-WMRC.369

**Purpose of study** The Mustarde otoplasty is a commonly used procedure for the correction of prominent ear. Complication rates following Mustarde otoplasty, related to suture extrusion and long-term outcomes, are variable in the literature. Here, we examine the efficacy and safety of the Mustarde otoplasty.

**Methods used** Retrospective data was collected on patients under 18 years-old who underwent primary otoplasty by the senior author between May 2009 – August 2018. Patient demographics, clinical presentation, intraoperative details, complications, follow-up, and patient/family

9JIM 2020:68(Suppl 1):A1–A228 A159

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Abstracts

A COMPLEX INTERPROFESSIONAL SIMULATION SCENARIO OF NECK MASS WITH CRITICAL AIRWAY OBSTRUCTION

1BT Ward*, 1T Meyer, 1M Montegut, 2K Roth, 1M Sardesai. 1University of Washington, Seattle, WA; 2Western University, London, ON, Canada

Purpose of study Emergency airway management is a low-frequency, high-acuity situation that all otolaryngology residents encounter. This requires situational awareness, sound decision-making, and strong interprofessional communication. Few curricula target these non-technical skills. Simulation can enhance technical and non-technical skills in a safe setting. We aimed to develop a complex simulation scenario that promotes non-technical skills among junior otolaryngology trainees.

Methods used A script of a scenario describing impending airway obstruction was developed by a multidisciplinary team from a deidentified patient case edited using iterative input to simplify and ensure generalizability. The scenario was tested on junior otolaryngology trainees using a high-fidelity mannequin in a simulated OR, appropriate instrumentation, anesthesiologist and nurse confederates, and a facilitator. In the scenario, a 52-year-old with a neck mass had worsening dyspnea that progressed to respiratory distress requiring intubation. The median operative time was 95 minutes (31-133 minutes). A total of 24 complications were reported in 17 patients. Complications included: Suture extrusion (n=20), hematoma (n=1), suture abscess (n=1) and reoperation (n=2). The study had a revision rate of 1.7% (n=2). No additional procedures were documented at other hospitals in the province. The majority (97%) of reported ear outcomes demonstrated both patient and surgeon satisfaction. The scenario was felt to be realistic by trainees and faculty, offered educational value across ACGME core competencies, and improved trainee confidence. The scenario may be a resource for programs implementing similar training.

Summary of results 18 R1 to R3 level trainees participated in the scenario. Trainees showed an increase in confidence with airway management from 2.3/5 to 3.1/5 which was not statistically significant. Participants scored the scenario realism as 4.18/5 and experts reviewing the case scored the case as 4.5/5. Seven expert evaluations showed agreement to strong agreement that the scenario enhanced medical knowledge (4.6), technical skill (4.4), communication (5.0), professionalism (4.7), and systems-based practice (4.6).

Conclusions This scenario was felt to be realistic by trainees and faculty, offered educational value across ACGME core competencies, and improved trainee confidence. The scenario may be a resource for programs implementing similar training.

371 DUAL-PLANE VERSUS PREPECTORAL BREAST RECONSTRUCTION IN HIGH BMI PATIENTS

1RM Gold*, 1E O’Rorke, 2M Pfaffenberger, 3L Eldenburg, 1T Rorke, 1A Gabriel. 1Elson S. Floyd College of Medicine, Spokane, WA; 2Private Practice Allen Gabriel MD, Vancouver, WA; 3Compass Oncology, Vancouver, WA; 4Loma Linda University Medical Center, Loma Linda, CA

Purpose of study Patients with high body mass index (BMI $\geq 30$ kg/m$^2$) who undergo breast reconstruction face unique challenges and postoperative complications associated with their BMI. The optimal reconstructive approach for these patients remains to be determined. In order to determine if there was an association between postoperative complications and the plane of reconstruction, this study compared outcomes of prepectoral and dual-plane reconstruction specifically in patients with high BMI.

Methods used This study included high BMI patients who underwent immediate dual-plane or prepectoral expander/implant reconstruction. Patients were stratified by reconstruction plane (dual-plane or prepectoral). Subsequently, postoperative complications were compared between the two groups. Multivariate logistic regression analysis was performed to determine whether the reconstruction plane was an independent predictor of any postoperative complication after adjusting for potential confounding differences in patient variables between the groups.

Summary of results 133 patients enrolled in the study. Of those, 63 (128 breasts) underwent dual-plane and 68 (129 breasts) underwent prepectoral reconstruction. Rates of seroma (13.3% vs 3.1%), surgical site infection (9.4% vs 2.3%), capsular contracture (7.0% vs 0.8%), and any complication (25.8% vs 14.7%) were significantly higher in patients who had dual-plane vs prepectoral reconstruction ($p<0.05$). Multivariate logistic regression identified the following variables as significant, independent predictors of any complication: dual-plane reconstruction, diabetes, neoadjuvant radiotherapy, and adjuvant chemotherapy ($p<0.05$). Dual plane increased the odds of any complication by 3-fold compared with the prepectoral plane.

Conclusions Compared with the dual-plane approach of reconstruction, this study demonstrated that the prepectoral approach was associated with a lower risk of postoperative complications following immediate expander/implant breast reconstruction and may be a better reconstructive option for high BMI patients.
BOTH THE GUIDEWIRE AND STENT SELECTED INFLUENCE THE ABILITY TO BYPASS AN IMPACTED URETERAL STONE
J Hutton*, AS Amayali, J Groegler, J Maldonado, R Belay, L Burke, N Brown, D Baldwin. Loma Linda University, Loma Linda, CA

10.1136/jim-2019-WMRC.372

Purpose of study This bench-top study compares the ability of 24 different guidewire-stent combinations to bypass an impacted ureteral stone.

Methods used A bench-top impacted ureteral stone model was used to compare the force required to bypass a tightly impacted 5 mm ureteral Begostone. 6 different wires and 4 different stents were placed beyond an impacted stone using a Mark-10 Force Gauge. 10 trials were run for each stent-guidewire combination unless no stent passage occurred after 5 trials (N=190). Primary outcomes included stent passage rate and insertion force (lbf). The stent passage rates were compared using a Chi-Square test. The average stent insertion force was compared using ANOVA and Tukey tests. A p-value <0.05 was considered significant.

Summary of results The highest stent passage rates were seen in the Inlay-ZIPwire (100%) and Lubriflex-Glidewire (100%) (table 1-E). The stent passage rates were significantly different between the different combinations (Chi-Square test; p<0.05). The stent passage rate of each individual stent regardless of the guidewire used were significantly different (Chi-Square Test; p<0.05). The Olympus Lubriflex stent had the highest stent passage rate (78.18%; N=55), followed by the Bard Inlay Stent (63.64%; N=55). Of the stents that passed, the Inlay-ZIPwire required the least insertion force (table 1).

Conclusions These results suggest that both the guidewire and stent employed significantly influence the force required for stent passage. Knowledge of these factors could help surgeons select the optimal stent-guidewire combinations for these surgeries.

Abstract 372 Table 1 Stent passage rate and average insertion force required to bypass an impacted stone of 24 different stent-guidewire combinations in a benchtop model. Pearson Chi-Square test: p<0.05

<table>
<thead>
<tr>
<th>A. Stent</th>
<th>B. Guidewire Manufacturer</th>
<th>C. Guidewire</th>
<th>D. Stent Passage Rate</th>
<th>E. Average Insertion Force</th>
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<tbody>
<tr>
<td>Bard Inlay</td>
<td>Bard Boston Scientific</td>
<td>SOLO Hydro Hybrid</td>
<td>5/10 (50%)</td>
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<td>Cook Medical</td>
<td>ZIPwire</td>
<td>10/10 (100%)</td>
<td>0.21710</td>
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<td></td>
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<td>Amplatz Super Stiff</td>
<td>3/10 (30%)</td>
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<td></td>
<td>HiWire</td>
<td>8/10 (80%)</td>
<td>0.46300</td>
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<td>9/10 (90%)</td>
<td>0.30057</td>
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<td></td>
<td></td>
<td>Glidewire</td>
<td>0/5 (0%)</td>
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<td>Boston Scientific Percuflex</td>
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<td>ZIPwire</td>
<td>0/5 (0%)</td>
<td>N/A</td>
</tr>
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<td></td>
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<td></td>
<td>HiWire</td>
<td>0/5 (0%)</td>
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<td></td>
<td>UltraTrack</td>
<td>0/5 (0%)</td>
<td>N/A</td>
</tr>
<tr>
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<td>Olympus Lubriflex</td>
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A NEW HOUR-SPECIFIC SERUM BILIRUBIN NOMOGRAM CONSTRUCTED WITH DATA FROM 400,000 NEONATES

Methods used This study utilizes a retrospective cohort design to evaluate preterm infants born between May 2007 to July 2019. Children were assessed using the Bayley Scales of Infant and Toddler Development-Third Edition. Multivariable regression was used to evaluate the association between RBC transfusion and neurodevelopmental outcomes. Severe neurodevelopmental impairment (NDI) was defined as any Bayley composite score <70, blindness, deafness, or an abnormal neurological exam.

Summary of results 602 children were evaluated. Two hundred sixty-eight (45%) infants received a total of 691 RBC transfusion exposures. 52% received RBC transfusion after 72 hours of life. After adjusting for maternal age, gestational age, gender, intraventricular hemorrhage, and need for inotropes, RBC transfusion was associated with lower two-year cognitive, language, and motor composite scores (adjusted-β [95% CI]: -2.53 [-4.75 to -0.31], P=0.026; -2.64 [-5.25 to -0.03], P=0.047; -3.57 [-5.90 to -1.24], P=0.003, respectively). RBC transfusion was also associated with an increase in the adjusted-odds of developing severe NDI (adjusted-OR, 1.83 [95% CI: -1.02-3.30], P=0.043). In an adjusted analysis of preterm infants <28 weeks, RBC transfusion retained statistical significance for cognitive composite scores (P=0.038).

Conclusions RBC transfusion was an independent predictor of lower cognitive, language, and motor outcomes in preterm infants at two-years after adjustment for differences in perinatal characteristics and illness at birth. Prospective studies utilizing blood-saving strategies (placental transfusion, minimal blood draws), to reduce the need for blood transfusion and improve long term outcomes, are needed to confirm this association.

TELEHEALTH DIAGNOSES OF AUTISM SPECTRUM DISORDER IN TODDLERS

Purpose of study Determine telehealth accuracy and efficacy in diagnosing autism spectrum disorder in children aged 21 months to 48 months

Methods used Children receiving care at a military hospital in Alaska received comprehensive telehealth evaluations by two Developmental Pediatricians and fellows located at Madigan Army Medical Center, Joint Base Lewis Mcchord, WA; 2Bassett Army Community Hospital, Fairbanks, AK.

Summary of results Of the 15 children enrolled, 7 received Autism diagnoses after the telehealth evaluation (inter-rater reliability 100%). In-person evaluation diagnoses matched telehealth diagnoses in 15/15 children. 4/15 of the children underwent the confirmatory evaluation with a Developmental Pediatrician. The telehealth evaluation and confirmation evaluations were compared.

Conclusions This is the first study to report utilizing video telehealth to diagnose autism using a parent led play observation. This study supports the concept of conducting autism evaluations by video telehealth.
Behavior and development II
Concurrent session
8:00 AM
Saturday, January 25, 2020

376 MOTOR AND COGNITIVE DEVELOPMENTAL SCORES IN OFFSPRING OF PREGNANT SMOKERS AT 12 MONTHS OF AGE
T Batish*, K Miller, J Harris, A Vu, S Santon, C Morris, C McEvoy, Oregon Health and Sciences University, Portland, OR
10.1136/jim-2019-WMRC.376

Purpose of study In some studies, prenatal smoke exposure has been associated with decreased cognitive, motor, and behavioral function in the offspring. We conducted the current analysis to determine whether the neurodevelopmental scores collected by domain on the Ages and Stages Questionnaire (ASQ) scores at 12 months of age differed between offspring of prospectively identified pregnant smokers versus nonsmokers.

Methods used The validated ASQ-3 was obtained at 12 months of age for offspring delivered to pregnant smokers randomized to vitamin C versus placebo during pregnancy and offspring of pregnant nonsmokers also studied prospectively. Z-scores for each ASQ domain were calculated using the normative data obtained from the ASQ-3 user’s guide. The raw scores and Z-scores per domain were compared between groups using the Wilcoxon signed rank test.

Summary of results ASQ results were obtained in 204 offspring of smokers and 32 of nonsmokers. The demographics of the two groups were: Smokers: 19% non-white, 87% government assisted or self-paid/none insurance status, 54% ≤ high school education, 8 cigarettes/day at randomization into the study at <23 weeks of gestation. Nonsmokers: 6.3% non-white; 6.3% government insurance; 6.3% ≤ high school education. The offspring of smokers had significantly lower scores in the problem solving (Z score of -0.04 in smokers versus 0.32 in nonsmokers) and personal social (Z score of 0.04 in smokers versus 0.45 in nonsmokers) domains. See table 1.

Conclusions This data suggests a potential difference in problem solving and personal social domains in offspring of smokers versus nonsmokers. However, we are limited by our sample size to be able to adjust for other important covariates. Further study with longer follow-up and more detailed neurodevelopmental testing is needed.

Abstract 376 Table 1 Motor and cognitive scores in offspring of smokers and nonsmokers

<table>
<thead>
<tr>
<th>ASQ Domain</th>
<th>Offspring of Smokers (n=204)</th>
<th>Offspring of Nonsmokers (n=32)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Communication</td>
<td>51.3 ± 9.6</td>
<td>53.8 ± 8.5</td>
</tr>
<tr>
<td>Gross Motor</td>
<td>54.0 ± 10.2</td>
<td>53.1 ± 13.4</td>
</tr>
<tr>
<td>Fine Motor</td>
<td>54.6 ± 7.6</td>
<td>55.6 ± 6.6</td>
</tr>
<tr>
<td>Problem Solving</td>
<td>48.6 ± 10.4</td>
<td>52.5 ± 8.2*</td>
</tr>
<tr>
<td>Personal Social</td>
<td>46.2 ±12.5</td>
<td>51.1 ± 9.7*</td>
</tr>
</tbody>
</table>

377 THE IMPACT OF TELE-BEHAVIORAL HEALTH IN RURAL AMERICAN INDIAN COMMUNITIES
S Yang*, MK Lieng, J Mouzoon, I Sigal, JP Marcin, B Pakyurek, B Restrepo, JS Hoch, PM Yellowlees, University of California Health System, Sacramento, CA
10.1136/jim-2019-WMRC.377

Purpose of study Past studies on tele-behavioral health show that virtual psychiatric, substance abuse, and other mental health services are linked to improved quality of life and symptoms across demographically and diagnostically diverse groups of patients. However, there are still large geographically isolated populations, including American Indians, without access to behavioral health services and even less data demonstrating the effectiveness of tele-behavioral health services among these marginalized populations. The purpose of this study is to evaluate the impact of tele-behavioral health services provided to rural American Indian populations on patient-centered measures among both pediatric and adult patients.

Methods used Using our existing telehealth network with five Indian Health Services clinics located in rural American Indian tribal areas, we administered mental health surveys during patient-intake using five validated clinician-rated instruments: PHQ-9, GAD-7, PROMIS, DUDIT-C, CBCL; and two provider-observed validated instruments: CGI and GAF. For patients with more than one visit, the first two visits were compared using paired t-tests in R.

Summary of results Preliminary data from the past five months show a total of 122 completed tele-behavioral health appointments. From those, 17 self-reported mental health surveys and 59 provider-observed measures were completed. The baseline characteristics abstracted shows poor to moderate mental health symptoms for all patients across all validated instruments, with the exception of the drug usage disorder test, DUDIT-C, scoring low in usage. Furthermore, the data shows that for seven individuals who had more than one tele-health encounter over the 5 month period, there was a 4.42-point increase in GAF (95% CI: 1.10 to 7.76) from 60.3 to 64.7 which shows an improvement on the GAF scale from ‘Moderate symptoms’ to ‘Some mild symptoms.’

Conclusions Rural American Indian communities significantly suffer from mental health illnesses as well as barriers to care. The preliminary analysis of the short pre- and post- test evaluation indicate that tele-behavioral health services have the potential to improve mental health measures and illnesses in rural American Indian communities. Data collection will continue until March 2021.

378 PRIMARY CARE SCREENING PRACTICES FOR LEARNING DISABILITIES
1S Jonnalagadda*, 1R Boada, 2A Talmi. 1Children’s Hospital Colorado, Aurora, CO; 2University of Colorado School of Medicine, Aurora, CO
10.1136/jim-2019-WMRC.378

Purpose of study Regular screening can improve overall development through identification of children at risk for learning disabilities (LD) and facilitate more timely access to intervention. Failure to identify and address LD can result in academic failure, internalizing symptoms, and behavioral challenges. Currently, LD concerns are identified in the primary care setting when there are presenting concerns or through general developmental screens. We hypothesize that learning disabilities are
insufficiently identified using current primary care screening practices. This study aims to: a) characterize children who screen positive for learning concerns within our primary care clinic, b) compare percent of children who screen positive to prevalence rates of learning disabilities in the population and c) determine if a plan for intervention was documented by the primary care provider.

Methods used In the initial phase, we will retrospectively analyze electronic medical record data collected over an 18-month period. Children between ages 6 to 17 years with a positive screen for educational difficulties will be selected. Screens will be considered positive if the parent answers affirmatively to a question regarding school difficulties on the clinic’s psychosocial patient questionnaire. Data regarding other associated variables such as co-morbid conditions and socioeconomic status will also be obtained, to see how they correlate with positive LD screen status. Interventions in response to the positive screen will also be documented.

Summary of results Data collection and analysis is ongoing. Interim results including descriptive statistics of patients with positive screen will be presented.

Conclusions Results will be used to determine feasibility of integrating a specialized learning disability screener into current clinic workflow. We hypothesize that a specialized screener will improve identification of children at risk for learning disabilities when compared to current practice.

Developing Team NEMO: Restructuring the High-Risk Infant Follow Up (HRIF) Clinic for the Implementation of Evidence-Based Guidelines for Early Diagnosis of Cerebral Palsy (CP)

S Iyer, K Jiang*, S Espinosa, K Kesavan. UCLA, Los Angeles, CA

Purpose of study Recent guidelines recommend early diagnosis and early intervention to improve outcomes in children with CP. The HRIF clinic at University of California Los Angeles (UCLA) is part of a multi-site study aimed at implementing these guidelines and decreasing the age of CP diagnosis to under 12 months. Team NEMO (Neurodevelopment and Early Movements Observation) was formed to facilitate the implementation.

Methods used Setting: 2 HRIF clinics at UCLA.
Implementation stages:
- Exploration – leadership commitment, review of current practices, addressing barriers to implementation using SWOT analysis, IRB approval
- Installation – selecting staff, training, process flow design, redesign in clinic structure, schedule of visits
- Initial implementation – data collection, refining process flow design, weekly team meetings, monthly site calls
- Full implementation – monitoring fidelity of process flow, sustaining gains, making plans for further expansion to other sites

Patients who qualify for HRIF were identified during their stay in the neonatal intensive care unit (NICU). Babies who did not qualify for HRIF but with abnormal general movements assessment were also referred to the HRIF. Babies were seen in the HRIF clinic starting at 3-4 months corrected gestational age.

Summary of results Compared to the year before implementation, the mean age of CP diagnosis decreased from 15.4 months to 12.03 months. Total number of new diagnosis of CP increased from 4 to 8. The number of high-risk for CP diagnosis increased from 0 to 17, with average age for high risk of CP diagnosis being 3.7 months. The number of visits for babies under 4 months increased from 45 to 134. No-show rate decreased from 20% to 13%

Conclusions The NEMO program successfully implemented the guidelines, resulting in decrease in average age of cerebral palsy diagnosis to 12 months.

Primary Care Management of Attention-Deficit/Hyperactivity Disorder in Children Aged 2 – 5 Years

Y Bennett*, HM Feldman, RM Gardner, O Blaha, LC Huffman. Stanford University, Stanford, CA

Purpose of study Attention-deficit/hyperactivity disorder (ADHD) affects approximately 2% of US preschool-age children. Little is known regarding preschool ADHD diagnosis and management by primary care providers (PCPs). This study assessed (1) rates of PCP diagnosis of ADHD in preschool-age children, (2) PCP adherence to select aspects of ADHD clinical practice guidelines, and (3) patient factors influencing variation in PCP diagnosis and management.

Methods used Retrospective cohort study of electronic health record (EHR) data from all office visits of children aged 2-5 years, seen ≥2 times between 2015 and 2019, in 10 practices of a community-based California primary healthcare network. Study outcomes included ADHD diagnosis (symptom- or disorder-level), and adherence to guidelines in (1) documentation of comorbidities at or after an ADHD diagnosis, (2) choice of ADHD medication, and (3) follow-up of medicated patients.

Summary of results Of 29,408 eligible children aged 2 to 5 years, 195 (0.7%) carried an ADHD diagnosis (0.04%-0.85% from 2-5 years, respectively). Of those, 109 (56%) had only symptom-level ADHD diagnoses (e.g., hyperactivity); 105 (54%) had documented comorbidities (e.g., language delay/disorder). ADHD medications were prescribed only to children aged 4-5 years (40/195 (21%)); 34 of those 40 received stimulants as the first-line medication, and 19 of 40 had a follow-up visit within 2 months. In logistic regression models, children with public or military insurance were more likely to have an ADHD diagnosis (OR=1.64; CI: (1.17, 2.26); OR=2.22; CI: (1.32, 3.56)); ADHD patients initially diagnosed at a younger age and those with military insurance were more likely to have documented comorbidities (OR=1.06 (1.03, 1.10); OR=2.94; CI: (1.064, 5.29)).

Conclusions The rates of PCP diagnosis of ADHD in preschoolers in this network were below estimated population prevalence. PCPs mostly followed guidelines in identification of comorbid conditions and in choice of stimulant medications, but had low rates of timely follow-up. Preschool-age children with ADHD may have remained undetected, with evidence of sociodemographic disparities in diagnosis of young children with ADHD.
381 EARLY CHILDHOOD INTERVENTION SERVICES IN SEX CHROMOSOME ANEUPLOIDIES (SCA)

N Tartaglia, S Howell, S Davis, TG Thompson*. University of Colorado, Aurora, CO
10.1136/jim-2019-WMRC.381

Purpose of study Children with SCA such as XXY, XYY, XXX, and XYYY have higher rates of early developmental delays and many require special education supports. However, research on early childhood outcomes in SCA is limited and was conducted prior to current special education law and practices. It is critical to update our understanding of current community and educational supports for children with SCA in order to guide families and providers, and to inform early childhood systems and policy. This study examines parent-reported publicly funded intervention services in young children with SCA and parent perception of early childhood providers’ knowledge of SCA conditions.

Methods used Data were collected from a larger electronic REDCap survey (N=550) examining community and educational supports in parents of children 0-21 with a confirmed SCA diagnosis. Participants were recruited from clinics and advocacy organizations. Questions focused on participation in early intervention/birth to 2 services (EI) and early childhood special education.preschool (ECSE) and took 5-20 minutes to complete. The current subset (N=105) included US children who had not yet entered Kindergarten.

Summary of results Results show that over half of all children receive some kind of EI or ESCE therapy (EI=59%, ECSE=38.5%) and that tetrasomy SCA conditions (XXYY, XXXX, XXXXX) (EI=87%, ECSE=100%) are more likely to receive therapies than trisomy conditions (XXY, XYY, XXX)(EI=54%, ECSE=48%(p=0.023). Speech therapy is the most common therapy for both EI and ECSE groups across all SCA conditions. Most children receive 1 hour or less of EI therapies per week (96%) and the majority (72%) of preschool students are in inclusive classroom settings. Over half (57%) of parents report their child’s early childhood therapist and/or teachers have little to no knowledge of SCA conditions, and none(0%) were rated as ‘highly informed.’

Conclusions Medical and developmental providers should be aware of the frequent need for EI in infants and young children with SCA so that appropriate evaluations and interventions can be implemented in a timely manner. Parent report on the limited knowledge of SCA in early childhood providers and teachers justifies a need to train therapists, teachers, and policy makers in the unique developmental needs in children with SCA.

382 ETHNIC DISPARITIES IN REFERRALS AND SERVICES RECEIVED AFTER NICU FOLLOW-UP EVALUATIONS IN A SINGLE CENTER IN CALIFORNIA

RD Rodriguez*, UC Davis MIND Institute, Sacramento, CA
10.1136/jim-2019-WMRC.382

Purpose of study The NICU population is at an increased risk for neurodevelopmental impairments. Many of them are not being appropriately referred to Early Intervention (EI) for reasons that have not been fully explored. Families of low SES or minority groups may have more difficulty accessing these services. We hypothesized that 1-Non-White ethnicities have lower EI referral rates from High Risk Infant Follow-up (HRIF) clinics compared to Caucasians and 2-of those referred to EI, Non-White children have the lowest receipt of EI services compared to Caucasians.

Methods used This was a secondary analysis of data collected from HRIF visits through the California Perinatal Quality Care Collaborative (CPQCC) database at UC Davis from 2010-2014. Eligible patients were those who were evaluated at ≥18months and had a documented subsequent visit for which EI referrals were warranted.

Summary of results 135 individuals met inclusion criteria. Across individuals, the average age was 20.3 months at the 1st visit (SD=2.6, range=18-29). More Non-Whites were referred for speech (p=0.003) and developmental (p=0.03) services than Caucasians. A smaller percentage of Non-Whites were receiving PT (p=0.04), OT (p=0.04) and vision (p=0.01) services at the 1st visit. At the subsequent visit, barely half of those referred for speech therapy, and less than one third of those referred for other services (PT, Behavior, OT) were receiving appropriate services.

Conclusions Part C of the IDEA is intended to enhance the development of infants and toddlers with disabilities, minimize the need for special education and maximize the potential for independent living. Contrary to our hypotheses, we found that Non-Whites are being appropriately referred; however, further investigation into the reasons why Non-White children are not receiving appropriate services is warranted, such as individual barriers (medical complexity, language, culture, finances and family priorities, etc.) or systems issues (geography, wait list and capacity, budgets, eligibility differences). There is an urgent need to identify, document and understand the reasons for these disparities in services and develop targeted interventions to optimize development in this population.

383 VISION LOSS AND PSYCHOPATHOLOGY

JR Mark*, AG Kulinch, LM Scher, MJ Mannis. UC Davis School of Medicine, Sacramento, CA
10.1136/jim-2019-WMRC.383

Purpose of study By examining the associations between vision loss and various types of psychopathology within the current literature, this manuscript will provide ophthalmologists, psychiatrists, and psychologists insight into the relationships between vision and psychopathology to foster cross-professional relations. Topics explored include mental health consequences of vision loss, functional neurological disorders and deception syndromes, eye specific personality changes, Charles Bonnet syndrome, and vision disorders associated with psychosis.

Methods used A narrative literature review was conducted using PubMed, Google Scholar, and Scopus internet search engines. Study types examined included reviews, case studies, survey analyses, population-based studies, and twin studies among others.
Abstracts

385 EXCESS IODINE EXPOSURE: A CONTINUED RISK FOR SUBCLINICAL HYPOTHYROIDISM IN INFANTS

1C Kapoor*, 2A Jiang, 2JD Brandt. 1University of California, Davis, School of Medicine, Sacramento, CA; 2,3B Afghani. 1UC San Diego, La Jolla, CA; 2UC Irvine, School of Medicine, Irvine, CA; 3CHOC Hospital of Orange County, Orange, CA

Purpose of study The fetus and newborn infants are sensitive to the thyroid suppressive effects of excess iodine. The purpose of this study was to see whether exposure to excess iodine during pregnancy or lactation continues to be a risk to the newborn infants.

Methods used A literature search was utilized through search engines, PubMed, Google Scholar, and Cochrane using keywords, ‘iodine’, ‘neonate’, and ‘thyroid’. Studies included must have measured iodine concentration in urine and/or breast milk as well as thyroid function tests in the newborns or infants. Case reports were excluded.

Summary of results 4 studies met our inclusion criteria. All of the studies took place outside of the United States. Sources of excess iodine included water, household salt and seaweed products. Most studies showed an association between iodine excess, measured in urine or breast milk and the development of subclinical hypothyroidism. Because of lack of long term follow-up, it is unclear if the abnormal thyroid function tests persisted. Other limitations included lack of baseline sampling in some studies and lack of controls that would take into account other possible sources of excess iodine or other factors that could lead to abnormal thyroid tests.

Conclusions Our literature review suggests that excess iodine intake in certain countries worldwide continues to pose a health risk to newborns. Iodine intake as measured in the breastmilk or urine is associated with development of subclinical hypothyroidism. Healthcare professionals should be aware of socio-economic and cultural factors that could lead to intake of excess iodine. To minimize the risk to the newborn, pregnant mothers should be educated about the sources of excess iodine.

Clinical epidemiology and health disparities
Concurrent session
8:00 AM
Saturday, January 25, 2020

384 OUTCOMES IN PEDIATRIC GLAUCOMA

1C Kapoor*, 2A Jiang, 3JD Brandt. 1University of California, Davis, School of Medicine, Sacramento, CA; 2UC Davis Eye Center, University of California, Davis, Sacramento, CA

Purpose of study To determine visual outcomes and risk factors for vision loss for patients with childhood-onset glaucoma, to externally validate a provisional glaucoma severity scale, and to explore potential additional risk factors impact long term visual outcomes.

Methods used Retrospective review of electronic medical records of patients with a history of pediatric glaucoma. Data collected included initial presentation characteristics such as age, IOP, and refraction status, prior surgeries, acquired and non-acquired ocular conditions, glaucoma subtype, number and types of glaucoma surgeries, unilateral and bilateral disease, insurance type, patient adherence to prescribed follow-up (number of no-show appointments), along with demographic characteristics. Univariate analysis was used to determine associations between each putative factor and ultimate visual outcome.

Summary of results 177 patients with a diagnosis of childhood-onset glaucoma were identified; 153 eyes of 93 patients met inclusion criteria. Male:Female ratio was 49:51. Mean age at presentation was 4.69 ± 6.23 years. Mean duration of follow-up was 9.64 ± 3.60 years. Ethnicity (non-Hispanic and Hispanic or Latino) was associated with poor visual outcomes (p=0.005). Of our 93 patients, 52 (56%) had private insurance, 33 (35%) had Medicaid, and 8 (9%) had unknown insurance type; private insurance was associated with a better outcome (p=0.021). Many patients missed appointments (3.98 ± 5.44 average no-show rate) but increased no-shows were associated with poor visual acuity (p=0.024).

Conclusions Previously-documented risk factors for poor visual outcome include glaucoma subtype and ethnicity. We identified new risk factors including increased no-show appointments and non-private insurance type as strongly correlated with poor visual outcomes. These factors’ influence on visual outcomes can help develop targeted interventions for glaucoma care.
ENCOUNTERS WITH EMERGENCY MEDICAL SERVICES PRIOR TO FATAL OVERDOSE: AN OPPORTUNITY TO INTERVENE?

A Rollins, L Barnard, M Sadinle, RJ Haruff, CR Counts, RT Rea, J Hood. University of Washington School of Medicine, Seattle, WA; Public Health Seattle and King County, Seattle, WA; King County Emergency Medical Services, Seattle, WA

Purpose of study Although Emergency Medical Services (EMS) is often involved in acute lifesaving measures in overdose situations, little is known about the nature of EMS care provided to persons prior to fatal overdose. We evaluated the nature and frequency of EMS encounters with persons who suffered overdose death in the year prior to death.

Methods used We conducted a retrospective cohort study of all residents who had fatal overdose in King County, Washington in 2018 to determine if they had an EMS encounter during the year prior to death. Information was abstracted from EMS and medical examiner records. Death records were matched to EMS records using probabilistic linking followed by manual review validation; T-tests were used for statistical analysis.

Summary of results During 2018, there were 387 deaths due to overdose, 69% (N=266) involved an opioid and 54% involved a stimulant. Among the 387 decedents, 39.5% (n=153) had one or more EMS encounters in the study interval distinct from their death event, representing a total of 710 EMS encounters. Demographic characteristics were similar between those who did and did not have an EMS encounter (66% vs. 71% male, 77% vs. 80% white race, average age 48 vs. 46 years respectively). However, a large proportion with an EMS encounter were homeless (29% vs 9%, p=0.002). Among those with an encounter, 37% (n=56) had an encounter within a month and 58% (n=89) within 3 months of their death. Sixty percent (n=91) had one or more EMS encounters in the year prior to their death. The most common reason for an EMS encounter was drug(s) or alcohol which occurred in 52% of encounters, followed by injury/trauma (12%) and behavioral or psychiatric complaints (7%). Most encounters (81%) were transported to the hospital.

Conclusions Over one third (40%) of drug-overdose decedents had one or more EMS encounters in the year prior to their death. EMS may be an efficient strategy to identify those at high risk of overdose death. Future studies should evaluate
other characteristics of persons who have multiple encounters with EMS but do not die from overdose.

EVALUATION OF DEPRESSION AND ANXIETY IN WOMEN WITH POLYCYSTIC OVARY SYNDROME BY PHYSICIAN TRAINEES
Sh Pakhdikian*, Y Fernandez-Sweeney, F Dong, AL Nelson. Western University of Health Sciences, Pomona, CA
10.1136/jim-2019-WMRC.387

Purpose of study Polycystic ovary syndrome (PCOS) is the most common endocrinopathy that affects reproductive-aged women. Estimates are that up to 40% of women with PCOS have depression and/or anxiety, which far exceeds prevalence in the control population. Although not used as criteria to formally diagnose PCOS, it is important to assess patients for these associated conditions. The aim of this study was to survey resident physicians in California in select specialties, known to work with women, to understand their screening and diagnosis practices for PCOS, while also exploring their attitudes about mental health.

Methods used We surveyed PGY ≥ 2 residents in Family Medicine, Internal Medicine, Obstetrics and Gynecology, and fellows in Endocrinology and Reproductive Endocrinology. The questionnaire asked about demographics, patient population, screening methods, and opinions on mental health-related statements. 22 individuals initiated the survey, 13 completed it in its entirety.

Summary of results All participants reported their care for women with PCOS. 57% of resident physicians said they would routinely screen women for post-partum depression. However, nearly 42% rarely assessed women for mental health risks or only screened if there was evident history. On average, physicians were neutral (on a 5-point Likert Scale) when asked if they found it easy to talk to their patients about mental health issues. They also collectively disagreed on their ability to provide options for referral to mental health services in their practice.

Conclusions With the growth and impact of research on behavioral medicine, we have seen an increase in confidence in discussing mental health. However, based on our findings resident physicians do not feel skilled in their ability to provide these services. Future studies should aim to enhance the structure of the clinical interview to include mental health risks and assessments to better determine the correlation of mental health consequences of this diagnosis in the full range of patient populations.

DOES INTENTION TO BREASTFEED AMONG EXCLUSIVE MARIJUANA USING MOTHERS DIFFER COMPARED WITH OTHER SUBSTANCE USERS?
1D Wanasighe*, 2S Shah, 3A Bamanikar, 4D Aboudi, 5S Mikkilineni, 6O Shyong, 7N Wu, 8C Giblin, 9T Lavan, 10H Brumberg. 1Westchester Medical Center, Valhalla, NY; 2Jersey Shore University Medical Center, Neptune, NJ; 3Department of Health, New York, NY; 4School of Public Health, Valhalla, NY; 5Lake Erie College of Medicine, Erie, PA; 6Touro Medical College, New York, NY; 7Hamilton College, Clinton, NY
10.1136/jim-2019-WMRC.388

Purpose of study Coincident with increased legalization, marijuana (MJ) use is increasing among pregnant women and women of reproductive age. However, mothers’ intention to breastfeed (BF) when using MJ around the time of pregnancy is unclear. Therefore we sought to compare intention to BF among mothers exclusively using MJ with those using electronic cigarettes (ecig), tobacco products, multi-substances, and non-users.

Methods used This is an interim analysis of surveys from parents of live singletons merged with birth certificate data. Subjects were categorized into 5 groups: ecig use only (ECO), marijuana use only (MJO), tobacco products only (TPO), multi-substance users (MSU; ≥ 2 of, marijuana, illicit drugs, ecig & tobacco), & non-users (NU; neither ecig, tobacco, marijuana). Due to small numbers exclusive illicit drug users were not able to be analyzed separately. Intention to BF was compared across groups. Also assessed were adverse life events in the peri-conceptional period such as partner incarceration, job loss, & homelessness. Logistic regression controlled for race/ethnicity, age, education, & adverse life events.

Summary of results Of 935 mothers: 1% were ECO, 3% were MJO, 9% were TPO, 9% were MSU, 78% were NU. In bivariate analysis, NU were older & ECO & MJO were younger. NU were more likely to be educated, married & along with ECO-employed. MSU had the highest rates of adverse life events. NU had the highest rates of intention to BF (78%) & were significantly higher than TPO and MSU (p’s<0.001). MJO were just as resolved to BF as NU. In adjusted models, there was no difference in intention to BF between MJO and NU.

Conclusions Mothers using exclusive MJ are similarly committed to BF as non-users. As studies suggest providers focus on legal issues when counseling MJ mothers, our findings highlight the need for clinicians to discuss potential newborn exposure via BF in discussions with MJ women- both pre and postnatally.

RACE AND ETHNICITY MORE INFLUENTIAL THAN SOCIOECONOMIC STATUS FOR DETERMINING EXCLUSIVE BREASTFEEDING SUCCESS
A Dong*, VP Walker, T Grogan, KL Calkins. University of California Los Angeles, Los Angeles, CA
10.1136/jim-2019-WMRC.389

Purpose of study The American Academy of Pediatrics recommends exclusive breastfeeding for the first six months of life. Less than 25% of families meet this goal. This study examined predictors of exclusive breastfeeding success.

Methods used This prospective, secondary study analyzed breastfed (BF) and non-breastfed (nBF) neonates (greater than 35 weeks gestational age) at nursery discharge and 30 days of age.

Summary of results At nursery discharge, the BF cohort’s (n=346) mean birthweight of 3.4±0.4kg was greater than the nBF cohort (n=43) mean birthweight of 3.3±0.5kg (p<0.004). There were more vaginal deliveries (76% vs. 56%, p<0.006) and non-Hispanic fathers in the BF cohort vs. nBF cohort (76% vs. 58%, p<0.008). While not statistically significant, there was a trend of more non-Hispanic mothers and White mothers in the BF cohort. At 30 days of age, the BF cohort (n=135) was more likely to have non-Hispanic fathers (85% vs. 70%, p<0.002), and White mothers (80% vs. 69%, p<0.07) than the nBF cohort (n=70). Non-Hispanic mothers...
were 2.8 times more likely to exclusively breastfeed than mothers who were Hispanic, after adjusting for mode of delivery, birth weight, and socioeconomic status (SES) at 30 days (odds ratio 2.8 (95% CI 1.1-6.9), \( p=0.03 \)). Infants with Non-Hispanic fathers were 2.7 times more likely to exclusively breastfeed than mothers whose partners’ were Hispanic, after adjusting for mode of delivery, birth weight, and SES at 30 days (odds ratio 2.7 (95% CI 1.1-6.3), \( p=0.03 \)). There was no difference in SES between groups at 30 days of age.

**Conclusions** This study supports ethnic/racial differences in exclusive breastfeeding rates. Paternal ethnicity may provide an additional indicator of breastfeeding disparities. Addressing disparities in exclusive breastfeeding rates may warrant additional investigations into culturally-inclusive, family-centered, and prenatally-initiated interventions.

**Abstract 390 Figure 1** Racial/ethnic differences in time to ED visit

**Abstract 390 Table 1** Utilization and demographics

<table>
<thead>
<tr>
<th>Interpretation Utilization, n(%)</th>
<th>All subjects (n=55)</th>
<th>Question 1 (n=17)</th>
<th>Question 2 (n=22)</th>
<th>Question 3 (n=16)</th>
<th>p-value</th>
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<tbody>
<tr>
<td>Interpreter Utilization, n(%)</td>
<td>44 (80)</td>
<td>14 (82)</td>
<td>14 (64)</td>
<td>16 (100)</td>
<td>0.02</td>
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<tr>
<td>Maternal age (yr), mean ± SD</td>
<td>33.2±5.9</td>
<td>31.5±6.4</td>
<td>33.7±5.8</td>
<td>34.4±5.3</td>
<td>0.34</td>
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<td>Parity, mean ± SD</td>
<td>3.1±1.5</td>
<td>3.3±1.4</td>
<td>2.9±1.5</td>
<td>3.2±1.6</td>
<td>0.70</td>
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<td>Vaginal delivery, n(%)</td>
<td>37 (69)</td>
<td>11 (65)</td>
<td>14 (66)</td>
<td>12 (75)</td>
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<td>Public insurance, n(%)</td>
<td>53 (98)</td>
<td>17 (100)</td>
<td>20 (95)</td>
<td>16 (100)</td>
<td>0.99</td>
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<tr>
<td>Gestational age (wk), mean ± SD</td>
<td>38.5±1.5</td>
<td>38.2±1.4</td>
<td>38.5±1.5</td>
<td>38.7±1.6</td>
<td>0.61</td>
</tr>
</tbody>
</table>

**Conclusions** There are racial/ethnic disparities in ED utilization among preterm infants over the first year of life, with Black and Hispanic infants more likely to visit the emergency room compared to White non-Hispanic infants.

**Abstract 391 Table 1** Utilization and demographics

<table>
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<td>31.5±6.4</td>
<td>33.7±5.8</td>
<td>34.4±5.3</td>
<td>0.34</td>
</tr>
<tr>
<td>Parity, mean ± SD</td>
<td>3.1±1.5</td>
<td>3.3±1.4</td>
<td>2.9±1.5</td>
<td>3.2±1.6</td>
<td>0.70</td>
</tr>
<tr>
<td>Vaginal delivery, n(%)</td>
<td>37 (69)</td>
<td>11 (65)</td>
<td>14 (66)</td>
<td>12 (75)</td>
<td>0.80</td>
</tr>
<tr>
<td>Public insurance, n(%)</td>
<td>53 (98)</td>
<td>17 (100)</td>
<td>20 (95)</td>
<td>16 (100)</td>
<td>0.99</td>
</tr>
<tr>
<td>Gestational age (wk), mean ± SD</td>
<td>38.5±1.5</td>
<td>38.2±1.4</td>
<td>38.5±1.5</td>
<td>38.7±1.6</td>
<td>0.61</td>
</tr>
</tbody>
</table>
(16/16), respectively (p=0.02). The most common reasons for declining an interpreter were a family member or friend who offered to interpret and self-reported English-proficiency.

**Conclusions** All subjects offered an interpreter with Question 3 accepted interpretation services. Although limited by sample size, the results highlight an important finding, that the way an interpreter is offered impacts utilization. By choosing the right words, we may temper the language barrier that exists for many families and ultimately decrease health care disparities. This further underscores the need for heightened focus on how to best communicate with LEP patients.

**Endocrinology and metabolism III**

**Concurrent session**

**8:00 AM**

**Saturday, January 25, 2020**

**393** **PROMISCUOUS NFkB ACTIVATION INDUCES PANCREATIC β CELL DEATH FOLLOWING GENOTOXIC STRESS**

S McKenna*, R de Dios, T Nguyen, J Sandoval, P Rozance, C Wright. University of Colorado, Aurora, CO

10.1136/jim-2019-WMRC.392

**Purpose of study** Intrauterine growth restriction (IUGR) affects 5-15% of gestations and is associated with an increased risk of developing Type II Diabetes Mellitus (T2DM). However, the mechanisms linking IUGR and T2DM are unknown. Although multifactorial, T2DM partially develops because of oxidative stress leading to genotoxic damage in pancreatic β cells. Recent *in vivo* studies have shown that NFkB inhibitory protein levels are chronically lower in IUGR pancreatic islets. Chronically reduced NFkB inhibitory protein levels in the islet may sensitize IUGR individuals to genotoxic insult and hasten β cell dysfunction in T2DM. In this study, we sought to determine whether lowering the NFkB activation threshold by decreasing total cellular inhibitory protein content could sensitize pancreatic β cells to genotoxic stress and diabetes development.

**Methods used** To assess the role of NFkB in the response to genotoxic stress, MIN6 immortalized murine insulinoma cells were exposed to streptozotocin(STZ,0.5-8mM, 0-24h), or were transfected with siRNA against the NFkB inhibitory protein IkBβ and exposed to STZ. Cell death following transfection and STZ exposure was evaluated by trypan blue exclusion. Expression of the NFkB target gene iNOS was measured by RT-qPCR. To assess the role of IkBβ in diabetes pathogenesis *in vivo*, adult(6-8 wk) male wild type(WT,C57Bl6/J) and IkBβ/-/- mice were exposed to multiple low-dose streptozotocin(MLDS,40 mg/kg IP) and blood glucose was measured weekly for 7 weeks following exposure to monitor for hyperglycemia and diabetes.

**Summary of results** STZ exposure induced dose-dependent cell death in MIN6 cells. Following transfection, cell death was increased compared to dose-matched control. SiRNA transfection and subsequent STZ exposure induced iNOS gene expression compared to STZ exposure alone. Taken together, these results demonstrate that decreased IkBβ sensitizes MIN6 cells to genotoxic stress. In vivo, IkBβ/-/- mice developed hyperglycemia and diabetes more rapidly than WT mice following MLDS exposure, indicating that lower inhibitory protein levels sensitize the pancreas to genotoxic stress and directly contribute to β cell dysfunction *in vivo*.

**Conclusions** Lower NFkB inhibitory protein levels sensitize pancreatic β cells to genotoxic stress and may help to explain the link between IUGR and T2DM.

**393** **PSYCHOLOGICAL TREATMENT RESOURCE USE IN PATIENTS WITH TYPE 1 DIABETES WHO ENDORSED DEPRESSIVE SYMPTOMS OR SUICIDAL IDEATION**

KR Stanek*, EM Youngkin, S Majidi. University of Colorado, Aurora, CO

10.1136/jim-2019-WMRC.393

**Purpose of study** Studies have shown higher depression rates in youth with type 1 diabetes (T1D) compared to the general population. Literature on suicidal ideation in T1D and how to provide appropriate follow-up for youth with depression or suicidal ideation is sparse. The objectives of this study were to evaluate the psychological resource use in adolescents with T1D after initial endorsement of depressive symptoms and/or suicidal ideation (SI), and to compare T1D and psychological characteristics between those who utilized resources versus those who did not.

**Methods used** Patients at the Barbara Davis Center who scored positive for depressive symptoms or endorsed SI using the Patient Health Questionnaire 9 (PHQ-9) were identified by retrospective chart review. At time of PHQ-9 screening, patients with positive scores met with a provider or psychologist and were provided psychological resources for follow-up. T1D characteristics and use of psychological resources before and after the initial screen were recorded. Chi-square, t-tests, and descriptive analyses were used for data analysis.

**Summary of results** Of 1376 screened, 200 patients were identified with positive depressive symptoms (n=169) and/or positive SI (n=101). Of those identified, 52% (n=104) utilized psychological resources; specifically, 30% therapy, 5% psychiatric medications, and 17% both. Those who utilized resources had a higher initial PHQ-9 score (13.6 vs. 12.0, p=0.02), were female (53.2% vs. 37.5%, p=0.03), and utilized a continuous glucose monitor less often (22.1% vs. 34.4%, p=0.01). There was no difference between groups in race, age, T1D duration, insulin pump use, average follow-up PHQ-9 score, SI at initial screen or follow-up, or HbA1c.

**Conclusions** Although depressive symptoms and SI is common in the pediatric T1D population, only half those who have a positive screen utilize the resources given, indicating additional education and follow-up are needed to help patients obtain the appropriate counseling needed. While no changes in glycemic management were seen in those who utilized resources, HbA1c changes are gradual; thus, a longer interval may be needed to determine if HbA1c does improve. Future studies should investigate how to improve utilization of resources in youth with depressive symptoms or SI, particularly in males.
Case report
Diabetes mellitus is becoming more common in older children but it’s extremely rare in infants. While type 1 and type 2 have a multifactorial etiology, neonatal diabetes is caused by single-gene mutations. This is a case of an infant with persistent hyperglycemia found to have a PDX1-gene mutation, which causes less than 1% of neonatal diabetes.

A 1-month-old boy born term weighing 2523 grams presented with a fever of 39.5°C. His bloodwork incidentally showed an hyperglycemia of 371mg/dL along with glucosuria and osmotic polyuria. Genetic testing revealed that he was heterozygous for a pathogenic variant in the PDX1 gene.

Neonatal diabetes usually presents before six months of age. PDX1 is just one of more than 20 known causative genes of monogenic diabetes. Early signs of diabetes mellitus like polyuria and polydipsia are easily missed in this age set. Consequently, most infants present with failure to thrive, severe dehydration and diabetic ketoacidosis. The course and treatment of neonatal diabetes depends on the genotype. Therefore, genetic testing is strongly advised for any infant presenting with diabetes less than one year of age.

Persistent hyperglycemia in an infant should draw concern for neonatal diabetes, especially in the context of a history of IUGR, glucosuria and osmotic polyuria. Genetic testing should be performed in these cases. The clinical course of neonatal diabetes depends on the genotype. In this case, the infant presented with persistent hyperglycemia requiring basal-bolus regimen, but the mild phenotype allowed the patient to be successfully weaned off insulin.

Sources:

Purpose of study
In high glucose environments such as diabetes, the enzyme aldose reductase (AR) acts to metabolize glucose to sorbitol which is thought to contribute to diabetic complications, including cataract, posterior capsular opacification, retinopathy, and neuropathy. It is therefore imperative for the research community to use a standardized model to understand AR-linked pathologies. In this study, we took advantage of CRISPR/Cas9 to create mutations in the AR gene that knocked out AR activity (ARKO), which may reduce the risk of diabetic complications. We hope to use these models for future work in understanding the aforementioned pathologies.

Methods used
Single guide RNAs (gRNAs) following CRISPR/Cas9 editing on exon3 of the AR gene and in vitro fertilization (IVF) was performed. Founders were characterized with DNA sequencing, protein western blot, and tissue histology.

Summary of results
DNA sequencing, protein western blot analysis, and tissue histology verified a 16 nucleotides deletion and 4 nucleotides insertion, respectively, in two founder lines of ARKO mice. Transmission of mutant alleles followed expected Mendelian patterns in pedigrees from both founder lines. Functional deletion of AR gene expression was shown in ARKO eye, heart, and kidney tissues where AR is more abundant in wild type mice. Furthermore, structural differences were not seen in these tissues when compared to wild type mice.

Conclusions
Two founder lines of ARKO mice were produced and characterized in our study. We believe that these lines can serve as an important base in tackling diabetic pathologies and understanding their mechanisms.
and 10 mg daily of ezetimibe was added. His LDL-C previously was 70 mg/dL. While the patient was on 40 mg of atorvastatin and 10 mg of ezetimibe, his LDL-C was lowered to 25 mg/dL after the above medical therapy. Due to a concern for inhibition of cortisol biosynthesis by statins, an ACTH stimulation test was done that showed a normal adrenal response. In addition, the patient had no signs nor symptoms of adrenal insufficiency.

This case illustrates that very low levels of LDL-C may be safe and do not inhibit cortisol response to ACTH stimulation test in T1DM patients. This may be important for T1DM patients with cardiovascular disease who would benefit from a low LDL-C.

**Impaired Catabolism of Calcitriol: A Rare Cause of Hypercalcemia and Calciphylaxis**

A 62-year-old male with multiple medical co-morbidities including chronic kidney disease (CKD) requiring dialysis presented to the hospital with altered mental status and hypercalcemia. Review of records revealed intermittent hypercalcemia dating back 17 years. The patient denied use of calcium supplements or calcitriol. Examination revealed a frail, cachectic gentleman with areas of dry gangrene on his hands consistent with calciphylaxis. Laboratory evaluation showed serum calcium ranging from 10.3-12.5 (NL 8.4-10.4 mg/dL), ionized calcium 1.34-1.40 (NL 1.13-1.32 mmol/L), phosphorus 4.5-6.1 (NL 2.5-4.5 mg/dL), serum creatinine 4.4 (NL 0.7-1.2 mg/dL), intact parathyroid hormone (PTH) 33 (NL 10–65 pg/mL), 25-hydroxyvitamin D 39 (NL 20 – 100 mg/dL) and 1,25 dihydroxyvitamin D of 82-115 (NL 18-72 pg/mL). SPEP, UPEP and PTH-RP were unremarkable. Hypercalcemia was attributed to an excess of 1,25-D. He was treated with pamidronate and had improvement in serum calcium. Unfortunately, the patient expired. Autopsy found no evidence of granulomatous disease or occult malignancy. Testing for abnormal metabolic clearance of vitamin D returned after his death showing an elevated 23 hydroxvitamin D to 24,25-dihydroxyvitamin D ratio (68.97, normal <25 ng/mL), indicating reduced activity of 24-hydroxylase. This enzyme catalyzes the hydroxylation of the biologically active form of vitamin D (1,25-D) and its precursor (25-hydroxyvitamin D) to inactive calcitroic acid for excretion. CYP24A1, the gene encoding the cytochrome P450 component of the mitochondrial 24-hydroxylase enzyme, is expressed in target tissues for vitamin D.

**Conclusions**

Disorders due to impaired 24-hydroxylase activity span a wide clinical range extending from life-threatening infantile hypercalcemia to chronic mild hypercalcemia. Circulating levels of active 1,25-D are dependent upon the activity of 1-alpha-hydroxylase (responsible for synthesis) and 24-hydroxylase (responsible for activation). This case demonstrates the crucial role of 24-hydroxylase activity, even in the setting of CKD. Defective 24-hydroxylase activity should be considered in the differential diagnosis for patients with chronic hypercalcemia and elevated 1,25-D levels.

**VCP Inhibitor CB-5083: A Potential Treatment for VCP Inclusion Body Myopathy**

C Cheng*, L Weiss, L Ta, T Tor, TA Do, V Kimonis. University of California, Irvine, Irvine, CA

Purpose of study Mutations in the Valosin-containing protein (VCP) cause inclusion body myopathy with early-onset Paget...
FURTHER DELINEATING A RARE, YET CLINICALLY PATIENTS WITH GENETIC VARIANTS OF UNKNOWN CAN and performed a retrospective chart review of our Methods used We reviewed the current literature for cases of Crouzon syndrome found to have molecularly confirmed tumors. Here we report a 13-year-old girl initially followed for Crouzon syndrome with acanthosis nigricans (CAN) and at age 11, extensive odontogenic tumors. At age 13, she was evaluated by genetics and molecular testing revealed the p.Ala391Glu variant in the FGFR3 gene consistent with the diagnosis of CAN. While she is monitored closely, her odontogenic tumors are so extensive she is unlikely to be a candidate for maxillary advancement or surgical resection.

Conclusions Due to phenotypic similarities with Crouzon syndrome, there may be delays in diagnosis or even underdiagnosis of CAN. Accurately distinguishing the two syndromes is crucial to enable proper genetic counseling and clinical surveillance, hopefully, allowing for early surgical intervention. Identifying odontogenic tumors on physical exam is difficult, thus we would advocate for 3D CT scan after a new diagnosis of CAN and periodically to evaluate for tumor development.

400 FURTHER DELINEATING A RARE, YET CLINICALLY SIGNIFICANT FEATURE OF CROUZON SYNDROME WITH ACANTHOSIS NIGRICANS: ODONTOGENIC TUMORS

R Gates*, C Hale, D Matalon. Stanford University, Stanford, CA
10.1136/jim-2019-WMRC.400

Purpose of study Distinguishing between Crouzon syndrome and Crouzon syndrome with acanthosis nigricans (CAN) is important as the two are genetically and clinically distinct syndromes. CAN is typically due to one recurrent pathogenic variant, p.Ala391Glu, in the FGFR3 gene, while Crouzon syndrome is due to variants in the FGFR2 gene. CAN is associated with similar facial features to Crouzon syndrome making diagnosis difficult. However, other features often present in CAN include choanal atresia, apnea, posterior fossa abnormalities, renal and skeletal anomalies, and rarely, odontogenic tumors. Here we report a 13-year-old girl initially followed for Crouzon syndrome found to have molecularly confirmed CAN with multiple odontogenic tumors.

Methods used We reviewed the current literature for cases of CAN and performed a retrospective chart review of our cohort of patients seen in the craniostenosis clinic at Stanford Children’s Hospital.

Summary of results We identified 38 cases of CAN in the literature. Of these cases, only three are reported with odontogenic tumors, and only one underwent successful resection. We report a case of a 13-year-old girl who had an abnormal head shape at 4 months of age. At 10 months of age, she was noted to have Chiari I malformation, hydrocephalus, and bicoronal craniostenosis; she was clinically diagnosed with Crouzon syndrome. Over time, she was noted to have obstructive sleep apnea, global developmental delay, acanthosis nigricans, and at age 11, extensive odontogenic tumors. At age 13, she was evaluated by genetics and molecular testing revealed the p.Ala391Glu variant in the FGFR3 gene consistent with the diagnosis of CAN. While she is monitored closely, her odontogenic tumors are so extensive she is unlikely to be a candidate for maxillary advancement or surgical resection.

Conclusions Due to phenotypic similarities with Crouzon syndrome, there may be delays in diagnosis or even underdiagnosis of CAN. Accurately distinguishing the two syndromes is crucial to enable proper genetic counseling and clinical surveillance, hopefully, allowing for early surgical intervention. Identifying odontogenic tumors on physical exam is difficult, thus we would advocate for 3D CT scan after a new diagnosis of CAN and periodically to evaluate for tumor development.

401 PATIENTS WITH GENETIC VARIANTS OF UNKNOWN SIGNIFICANCE FOR PRIMARY CILIARY DYSKINESIA: A CASE SERIES

DC Corpus*, PC Do, J Moua. UCSF-Fresno, Fresno, CA
10.1136/jim-2019-WMRC.401

Purpose of study Primary Ciliary Dyskinesia (PCD) is a rare autosomal recessive genetic disorder affecting ciliary structure and function and causing oto-sino-pulmonary disease. There are several diagnostic modalities to help confirm PCD including genetic testing.

Genetic testing for PCD is ever expanding, however the clinical significance of certain gene variants is not well established. Exploring these variants of unknown significance (VUS) may provide further insight into their clinical relevance. This case series seeks to characterize the clinical presentations and course of a subset patients with VUS for PCD.

Methods used Retrospective chart review of 11 cases were selected from patients of the UCSF Fresno Pediatric Pulmonology Clinic in Fresno, CA. Ten individuals had genetic testing performed due to symptoms and clinical history suspect for PCD. There was 1 case of PCD diagnosed via TEM. Diagnostic clinical predictors were included from the European Respiratory Society (PACDAR score) and American Thoracic Society criteria. Pulmonary function testings (PFT) and respiratory cultures were recorded as well.

Summary of results More than half (n=6) of the subjects were female, over the age of 13 years, and had PACDAR scores ≥4. A majority fulfilled at least 2 of the American Thoracic Society (ATS) criteria for PCD. All gene variants were associated with autosomal recessive disorders, primarily PCD, as well as Kartagener (n=1), infantile nephronophthisis (n=1), and Usher syndrome (n=1). Most patients had positive
Abstract 401 Table 1  Gene variants and clinical history of the sample

<table>
<thead>
<tr>
<th>Patient</th>
<th>PCD1</th>
<th>PCD2</th>
<th>PCD3</th>
<th>PCD4</th>
<th>PCD5</th>
<th>PCD6</th>
<th>PCD7</th>
<th>PCD8</th>
<th>PCD9</th>
<th>PCD10</th>
</tr>
</thead>
<tbody>
<tr>
<td>Clinical Summary</td>
<td>history of asthma, chronic rhinosinusitis, recurrent otitis media, pulmonary hypoplasia and dextrocardia</td>
<td>a history of asthma, chronic rhinosinusitis, recurrent otitis media, pulmonary hypoplasia and dextrocardia</td>
<td>history of multiple failed extubations, bowel perforation, CLD, interstitial emphysema, bronchiolitis obliterans vs BPD, ventriculomegaly</td>
<td>history of multiple failed extubations, bowel perforation, CLD, interstitial emphysema, bronchiolitis obliterans vs BPD, ventriculomegaly</td>
<td>recurrent sinus infections + cough occur every 2-3 months</td>
<td>recurrent severe respiratory infection and aspergillosis; otitis media, sinusitis, pneumonia</td>
<td>recurrent and chronic sinusitis, recurrent croup</td>
<td>recurrent and chronic sinusitis, recurrent croup</td>
<td>recurrent and chronic sinusitis, recurrent croup</td>
<td>recurrent and chronic sinusitis, recurrent croup</td>
</tr>
</tbody>
</table>

CLD: Chronic lung disease; BPD: bronchopulmonary dysplasia; **PCD 7 was the only patient with a 3rd gene variant

References

1. Versaci *, N. Dykeze, D. Chu, D. Stevenson. Stanford University, Stanford, CA; 2. Lucile Packard Children’s Hospital, Stanford, CA

Purpose of study Nevus comedonicus (NC) is a subtype of epi- dermal nevus originating from the hair follicle. It has been hypothesized that NC is due to somatic mutations in several candidate genes (e.g. FGFR2, NEK9). Levinsohn et al. identified somatic mutations in NEK9 in three cases of localized NC, each in highly conserved residues. NEK9 encodes a kinase important for centrosome separation. We hypothesize that the degree of mosaicism and tissue involvement of NEK9 leads to a broader NC syndrome than previously reported.

Methods used Chart review, physical examination, and literature review.

Summary of results Here we describe a 5 yo boy with leg length discrepancy, tibial dysplasia, metatarsus adductus, hemiparesis, tethered cord and agenesis of the corpus callosum with subcortical heterotopia. His skin exam was notable for many open comedones arranged in a Blaschkolinear distribution along bilateral axillae, chest and abdomen consistent with NC. The Blaschkoid pattern of the NC suggests a somatic mosaic event. In addition, extensive skeletal and nervous system findings suggest broader involvement. Molecular testing is scheduled on skin biopsy with paired blood for NEK9.

Conclusions We hypothesize that the phenotype of a mosaic mosaic is broad (e.g. PIK3CA-related conditions). A few isolated cases of NC are now reported to be secondary to mosaic NEK9 variants. Our case provides evidence of a broader NC syndrome with multigene involvement, which we hypothesize is due to somatic NEK9 variants in other tissue types, most often of the neurologic, ocular and musculoskeletal systems. Levinsohn et al. showed that mutations in NEK9 impair follicular cell differentiation in NC, although precise mechanisms in other ectoderm/mesoderm derived tissues has yet to be explored. The patient we present has significant systemic involvement; with unique features such as tibial dysplasia, metatarsal adductus, and cortical abnormalities. It is possible that similar to isolated NC, that other isolated phenotypes (e.g. isolated tibial dysplasia not due to NF1) could be secondary to somatic NEK9 variants.

403 MITOCHONDRIAL COMPLEX III DEFICIENCY IN AN ADOLESCENT WITH TRISOMY 21

C Vargas*, C Sosa. Valley Children’s Hospital, Madera, CA

Case report Mitochondrial Complex III Deficiency, nuclear type 2 (MC3DN2) is a rare autosomal recessive, severe progressive neurodegenerative disorder which results from a

A174 JIM 2020;68(Suppl 1):A1–A228
deficiency in the TTC19 gene. This encodes a respiratory chain protein that maintains the structural and functional integrity of the mitochondrial electron transport chain. There have been limited case reports of MC3DN2, to date 12 cases have been identified.

A 15-year-old male with Trisomy 21 consulted his pediatrician endorsing 3 weeks of toe-walking, ataxia and bilateral ankle contractures. One week later he developed dysphagia and dysphonia. On exam he had dysarthria, bilateral lower extremity hypertonia, hyperreflexia of the patellar tendons, wide based gait, pes cavus and positive Romberg sign. His complete blood count and metabolic panel, inflammatory markers, creatine kinase, and thyroid studies were normal. A lumbar puncture was normal and negative for oligoclonal bands, fungal, and bacterial cultures. A MRI of the head showed stable cerebellar atrophy and a right-sided brainstem lesion involving the olivary nucleus and the posterior medullary pyramidal. He was diagnosed with a demyelinating disease of unclear etiology. He received a 5-day course of high dose intravenous steroids and a repeat MRI showed no improvement. Upon follow up with his pediatrician, he was referred to genetics and mitochondrial sequencing was sent. Mitochondrial sequencing revealed a positive compound heterozygous mutation in the TTC19 gene and he was diagnosed with MC3DN2.

The symptoms seen in MC3DN2 are neurological, developmental, or behavioral in origin. The manifestations noted in this patient were dysphagia, aphasia, ataxia and muscle weakness. The patient’s neurologic status rapidly declined with loss of fine and gross motor skills, urinary and fecal incontinence, progression of dysphagia, loss of speech, and eventual death in a span of 9 months after diagnosis despite treatment with a mitochondrial cocktail. Studies show that there is a marked variation in rates of disease progression–progressive over decades to rapid progression to death.

This case illustrates the variable clinical presentation and rarity of MC3DN2 in the medical community; as well as its rapid progression to death in this patient with Trisomy 21.

404 SKELETAL DYSPLASIAS: A STANDARDIZED PROTOCOL FOR PRENATAL DIAGNOSIS

| 1JA Morales*, 2A Niemi, 3A Ness, 4D Krakow, 1L Hudgins. 1Stanford University, Stanford, CA; 2University of California, San Diego, La Jolla, CA; 3University of California, Los Angeles, Los Angeles, CA |

Purpose of study Develop an easily applicable standardized protocol for stepwise ultrasonographic evaluation of a fetus with short long bones suspected to have a skeletal dysplasia.

Methods used Prospective evaluation of fetuses suspected to have a skeletal dysplasia through a protocol developed at our institution based on a published guideline for the evaluation of a fetus with short long bones. All were re-evaluated following delivery or termination.

Summary of results Of 15 eligible fetuses, our protocol identified a probable diagnosis in 8 (53%) prenatally by ultrasound. In the 7 remaining, 3 had a ‘possible’ diagnosis (20%) and 4 were ‘unknown’ (27%). Postnatally, diagnosis was confirmed in all 8 probable cases: thanatophoric dysplasia (TD, n=4), osteogenesis imperfecta (OI, n=2), diastrophic dysplasia (1) and Ellis van Creveld (1). Of the 3 ‘possible’ diagnoses the following were observed: achondroplasia to unknown, Roberts phocomelia to TD, achondroplasia to Costello syndrome. In the ‘unknown’ category, 3 diagnoses were made postnatally: Kniest dysplasia, bent-bone dysplasia and mild OI. One ‘possible’ and one ‘unknown’ diagnosis were identified postnatally.

Abstract 404 Figure 1
remain as ‘unknown’, despite extensive clinical and radiographic review by experts.

Conclusions: We successfully identified the diagnosis by ultrasound in more than half (53%) of the cases. We observed that it was important to use the term ‘possible’ and ‘unknown’ when diagnosis was unclear, as the form of skeletal dysplasia remained uncertain postnatally. We believe that we can improve our detection rate by adding criteria for RASopathies and bent bone dysplasia as prenatal diagnostic criteria has been recently published for these conditions. We plan to perform a prospective analysis using our improved criteria.

Abstracts

405 NOVEL FINDINGS IN A PLCB4-RELATED DISORDER:
HYPERINSULINISM, VASOSPASM, AND CARDIAC
HYPERTROPHY

1A Kumar*, 1H Byers, 1H Hudgins, 2MA Deardoff, 1JA Bernstein. 1Stanford University, Palo Alto, CA; 2CHOP, Philadelphia, PA

10.1136/jim-2019-WMRC.405

Purpose of study: PLCB4 plays a role in endothelin signaling, regulating vascular tone in conjunction with nitric oxide and other factors. The gene is also implicated in auriculocondylar syndrome, an autosomal dominant condition characterized by ear and jaw anomalies. We describe two cases associated with the gene.

Methods used: Clinical phenotyping, chart review, whole exome sequencing and functional studies on patient-derived fibroblasts.

Summary of results: Patient 1 was born with macrosomia, hyperinsulinism, and cardiac hypertrophy. Postnatally, she developed marked digital and renal artery spasm as well as pulmonary hypertension requiring inhaled nitric oxide. Her appearance was notable for micrognathia, question mark ears and lower extremity melanocytosis. At 4.5 months she developed rapid onset of poor peripheral perfusion and passed away secondary to cardiac and respiratory decompensation. Trio exome sequencing revealed a de novo p.D630V variant in PLCB4. Patient 2 also exhibited hyperinsulinism and biventricular cardiac hypertrophy. She did not have jaw or ear anomalies but exhibited hypotonia and melanocytosis on her lower extremities. During infancy she exhibited abnormal movements. Brain MRI identified Moya-moya disease. She also experienced episodes of vasospasm of her peripheral vasculature. Exome sequencing showed a de novo p.D630N variant in PLCB4. She passed away at age 2 years due to cardiorespiratory failure after a neurosurgical procedure.

Conclusions: Somatic mutations p.D630V and p.D630N have been previously observed in uveal melanoma and are suspected to play a role in driving tumorigenesis. The fact that both individuals carry mutations in the same codon, separate from typical auriculocondylar hotspot codons 360 and 621 suggest a distinct mechanism of action.

Substitutions at codon p.D630V appear to be associated with vasospasm, cardiac hypertrophy and early death with or without typical features of auriculocondylar syndrome. Functional studies from patient-derived fibroblasts are underway; we hypothesize that these variants in PLCB4 may act via endothelin signaling to promote vasoconstriction.

Global health III
Concurrent session
8:00 AM
Saturday, January 25, 2020

406 LONG-TERM FEASIBILITY, SUSTAINABILITY, AND COST-BENEFIT ANALYSIS OF REUSABLE MENSTRUAL HYGIENE MANAGEMENT PRODUCTS IN A POPULATION OF ADOLESCENT FEMALES IN A REMOTE HIMALAYAN COMMUNITY


Purpose of study: Menstrual hygiene management (MHM) is often neglected in rural and remote communities due to many factors including: access, availability, education, sanitation, and cultural beliefs. This study aimed to better understand MHM practices and factors that influence them as well as assess the use, preference, and practicality of reusable pads and menstrual cups in a rural and remote school in the Indian Himalayas.

Methods used: In this cohort study, female students ≥13 years of age who have undergone menarche were recruited during annual health screens. In 2017, all participants were provided with locally made products: 1 menstrual cup and 4 reusable pads, and cleaning supplies. Product preference was then assessed over 24 months with individual interviews and questionnaires.

Summary of results: Of the 42 eligible females recruited, 100% chose to participate, 23.8% attended follow-up interviews and 48% completed follow-up questionnaires. In 2018, 80%, 30%, and 20% of participants preferred using the reusable pad, menstrual cup, and disposable pad, respectively. However, in 2019, product preference shifted to 42.9%, 9.5%, and 47.6%, respectively. Common reasons stated for discontinuation of reusable products included wearing-down of the product and having lost the product. Overall, a significant preference for the reusable pad compared to the menstrual cup was observed (p<0.05). Additional analysis of the perceived benefits and challenges of using reusable products was also assessed, including pain and discomfort (16.7% pad, 60.0% cup), fear (11.1% pad, 83.3% cup), leaks (21.2% pad, 66.7% cup), and difficulty cleaning (57.9% pad, 0.0% cup). While participants had significantly more fears about using the cup (p<0.01), they found the cup easier to clean than the reusable pads (p<0.05). Preliminary thematic analysis revealed that adequate menstrual blood containment, cost, and discreetness are important factors affecting MHM preferences among our participant population.

Conclusions: Overall, the findings suggest that reusable MHM products are feasible, culturally acceptable, cost-effective, and sustainable alternatives for women living in remote, resource-limited settings.
IMPROVED MATERNAL IODINE STATUS DURING FIRST TRIMESTER GESTATION WITH PRECONCEPTION NUTRITION SUPPLEMENTATION: THE WOMEN FIRST MATERNAL PRECONCEPTION NUTRITION TRIAL

A Young, J Kermo, C Uhtison, A Hendrick, J Westcott, S Ali, S Saleem, A Garroz, L Figueroa, N Somararav, S Goudar, M Hubmidge, B Kebs. Univ of CO School of Medicine, Denver, CO; 2Univ of CO Denver, Denver, CO; 3Aga Khan Univ, Karachi, Pakistan; 4Institute of Nutrition in Central America and Panama (INCAP), Guatemala City, Guatemala; 5KLE Academy of Higher Ed and Research's JN Medical College, Karnataka, India

Purpose of study Iodine deficiency remains common globally, with ~25% of low and middle-income country households without iodized salt. With its critical role in embryogenesis, iodine supplementation prior to conception may be important for fetal development. This study compared the effect of preconception nutrition supplementation (vs no supplementation) on maternal iodine status at 12 wk gestation and birth length in 3 low resource settings in the Women First Trial.

Methods used Women in Guatemala, India and Pakistan (n~100/arm/site) were randomized to receive multi-micronutrient fortified lipid-based nutrient supplements until delivery, starting ≥3 mo prior to conception (Arm 1), or starting at ~12 wk gestation (Arm 2) after sample collections. Urine iodine (µg)/creatinine (g) ratios (I/Cr) were determined at 12 wk. Cutoff for deficiency was I/Cr<1.50. Birth outcomes included length-for-age Z-score and rates of stunting (LAZ<-2) according to I/Cr status.

Summary of results At 12 wk mean ±SD I/Cr for Arm 1 vs Arm 2 was significantly higher for Guatemala (308±276 vs 200±137, p<0.0004) but not India (309±260 vs 243±306, p=0.09) or Pakistan (187±110 vs 205±132, p=0.30). Prevalence of deficiency was lower in Arm 1 vs Arm 2 in Guatemala (30% vs 44%, p=0.06) and India (24% vs 47%, p<0.0004); >40% were deficient for both arms in Pakistan. For combined sites and arms, means of birth LAZ were -0.71 ±0.94 and -0.59±1.07 (p=0.12) for I/Cr deficiency vs sufficiency ranges, respectively; newborn stunting rates did not differ by group.

Conclusions The preconception intervention resulted in improved iodine status during the first trimester in 2 of the 3 sites. The prevalence of iodine deficiency at 12 wk in all sites suggests opportunities for enhanced implementation of iodine fortification programs.

INCREASED DELIVERY SUPPORT FOLLOWING A SUSTAINABLE BIRTH CENTER PROGRAM IN A MOUNTAINOUS DISTRICT OF NEPAL

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Purpose of study Maternal and neonatal mortality have not improved in all regions of low resource countries with most of the recent progress occurring in urban areas. Health systems in the mountainous regions of Nepal have significant challenges in offering services. The neonatal mortality rate in the mountainous regions of Nepal are nearly double that of urban areas (45.6 vs 25.3 per 1,000 births).

This study aims to evaluate a sustainable birth center facilitation and birth attendant training program in a mountainous district of Nepal.

Methods used Solukhumbu District has a population of 100,000 people and is in a mountainous region of Nepal. A program was created with collaboration from the local community, the Ministry of Health, Nepali and US based NGOs, and the University of Utah. Pre-existing health posts in villages were equipped with delivery equipment. Health post staff were trained using the American Academy of Pediatrics and World Health Organization programs Helping Babies Survive and Helping Mothers Survive. The local community provides ongoing salary for the birth attendants and operating costs of the birth centers.

Women who delivered within the last two years were interviewed before and after implementation of the program. Interviews took place in 9 villages in Solukhumbu District between December 2015 and March 2019. Interviews were conducted in Nepali and data was analyzed using STATA.

Summary of results 363 women were interviewed prior to implementation and 243 women were interviewed following implementation. Prior to implementation 34.4% (125/363) of women interviewed delivered at a birth facility compared to 58.0% (141/243) following implementation [p<0.001, Chi square]. Prior to implementation 39.9% (144/361) of women interviewed delivered with a trained birth attendant present compared to 60.9% (148/243) following implementation [p<0.001, Chi square].

Conclusions Following implementation, there was an increased prevalence of women delivering at birth facilities supported by a trained birth attendant. This intervention shows promise in making quality birth health services more accessible to women and infants living in mountainous regions of Nepal. We hope to expand this pilot program into other mountainous districts as well as other low resource areas of the world.

HILLS BEYOND HILLS: AN ANALYSIS OF HEALTH CONDITIONS IN NORTHERN HILL TRIBES OF THAILAND

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Purpose of study The rural Hill Tribes of Northern Thailand are home to over fifty indigenous villages, and five main tribes. Inhabitants are refugees from Burma, Tibet, and China who migrated and settled in Northern Thailand’s forests 4,500 years ago. Between June 13th and June 20th, 2019, thirty-four medical students embarked on a medical mission trip with the Where There Is No Doctor non-profit organization, established and led by Dr. David Mar Naw. During the trip, students set up a medical clinic in two villages, serving the people of the Lahu tribe. Students became increasingly aware of the lack of available information about traveling to the Northern Hill Tribes. This project aims to provide an informational analysis of classic demographics, diagnoses, and treatments seen in the Northern Hill tribes, so that future medical providers traveling to this region can come prepared to provide better care.

Methods used Participants for this study were the Hill Tribe people of northern Thailand seeking medical care from healthcare providers of Where There is No Doctor organization. Patient information was protected through the use of a
random number generator. Healthcare providers first obtained subjective information through a questionnaire consisting of age, gender, past medical history, surgical history, social history and symptoms. Vital signs and objective information was then obtained using standard medical equipment. Patients were further questioned about chief complaint and associated symptoms with Dr. Naw, followed by administration of medications and a treatment plan.

**Summary of results** Results of the analysis showed a total of eighty-one participants, twenty-two of whom were pediatric patients (<18 years old), fifty-one were adult patients (aged 18 to 65), and five were geriatric patients (aged 65+). For the population seen as a whole, the most prevalent conditions observed were musculoskeletal pain (22 patients), poor appetite (20) and gastritis (18). The medications most frequently used were Omeprazole (19), Paracetamol (20), Vitamin B6 (24), and Analgesic Cream (22).

**Conclusions** With this clinical profile, future healthcare providers traveling to the Northern Hill tribes can be better prepared for their journey, and as a result, will make a lasting impact on the communities served.

### Abstracts

**410 ASSESSMENT OF THE SICKLE CELL DISEASE SCREENING PROGRAM IN THE INDIGENOUS THARU POPULATION IN NEPAL**

L Nakajima*, I Dascalu, K Marchand, A Khaira, S Yu, R Xu. University of British Columbia, Vancouver, BC, Canada

**Purpose of study** In 2015, UBC medical students, in collaboration with a Nepali community-based organization, CP Nepal, began screening for sickle cell disease (SCD) among the indigenous Tharu population in Dang, Nepal and estimated the prevalence of sickle cell trait to be 9.3%. SCD is an inherited disorder that causes painful vasoocclusive crises and significantly impacts quality of life. The Nepalese government has recognized SCD as a health crisis and funds treatment for those with a formal diagnosis of SCD. Since 2015, our team has worked with CP Nepal to equip local health posts with trained staff and laboratory capability to increase screening rates. This study examines the screening statistics collected since 2015 to evaluate the effectiveness of these initiatives.

**Methods used** Following the initial mass screening, 3 health posts continued providing screening in the community. From 2015-August 2019, data was collected by CP Nepal on screening rates and diagnostic test results of those that screened positive. Data was stored on a password encrypted excel spreadsheet.

**Summary of results** From 2015-2019, 4029 individuals were screened for SCD. In 2015, 2899 were screened (272 positive, 96 trait, 4 disease). In 2016, 144 were screened (24 positive, 5 trait, 2 disease). In 2017, 668 were screened (62 positive, 60 trait, 2 disease). In 2018, 114 were screened (14 positive, 12 trait, 1 disease). In 2019, 204 were screened (12 positive, 11 trait, 1 disease). Of those that got diagnostic testing, 184 individuals had sickle cell trait, and 10 had SCD.

**Conclusions** Organizing a mass screening appears to be the most effective in screening a high number of individuals. Although screening rates are variable, an increasing proportion of those that screen positive are getting further diagnostic testing. In 2019, 2 new health posts were established, which may explain the increased screening rates compared to 2018. Screening rates may be reaching a plateau due to saturation, thus future teams should expand efforts to surrounding regions. A limitation of this study is that the data is missing false positive screening rates and number of people getting treatment. Future studies should investigate the barriers that prevent those who screen positive from getting diagnostic testing and accessing treatment.

#### IMPROVING COMMUNITY AWARENESS OF SICKLE CELL DISEASE THROUGH EDUCATIONAL MODULES IN THE INDIGENOUS THARU POPULATION OF NEPAL


**Purpose of study** Sickle cell disease (SCD) is an inherited hemoglobinopathy which leads to increased morbidity and mortality and decreased quality of life. The prevalence of the sickled hemoglobin allele is estimated to be 9.3% among the indigenous Tharu population in Dang, Nepal. While the Nepalese government offers funding for those with a SCD diagnosis, a major barrier preventing access to care is lack of awareness about the disease and the screening process. Since 2015, our team has attempted to address this gap by delivering educational modules about SCD to communities in Dang, in hope of improving health literacy and encouraging individuals to get screened. This study examines the effectiveness of these modules by comparing participants’ understanding of the disease before and after their delivery.

**Methods used** Educational modules developed by our team covered SCD symptoms, pathophysiology, heritability, and the process of diagnosis and treatment in Nepal. Modules were delivered to community mothers’ groups in the local language by community health workers trained by our team. Binary response questionnaires assessing understanding of SCD and willingness to get screened were administered before and after the educational sessions. Participant responses were coded and analyzed using SPSS.

**Summary of results** A total of 81 mothers (mean age=36.1 years) participated in the study. Participants’ overall survey scores improved by 28.2% from pre- to post-educational questionnaires (mean score 51.9% pre-module (SD 11.9%), 80.1% post-module (SD 24.2%), P<0.001). Prior to the module, only 11.1% of participants reported having been screened. After the module, 86.8% of those not screened reported that they would get screened, and 88.2% would get their children screened.

**Conclusions** The data suggests that the educational modules were effective in improving participants’ knowledge about SCD and communicating the importance of getting screened, as the majority of participants reported willingness to get themselves and their children screened in the future. Further studies should investigate the impact of the educational modules on screening, diagnosis and treatment. Low literacy among the local population may limit the validity of the questionnaires.
HEALTH EDUCATION FOR PARENTS AND CAREGIVERS OF PEDIATRIC PATIENTS AT THE UGANDA CANCER INSTITUTE

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10.1136/jim-2019-WMRC.412

Purpose of study The Uganda Cancer Institute (UCI) sees patients with little understanding of cancer and low health literacy. There are widespread myths about cancer in Uganda, leading to late presentations requiring complex care. This project aimed to increase understanding of cancer and its treatment among parents and other caregivers at the UCI.

Methods used A community assessment and literature review on health education in low-literacy populations were conducted, and 5 UCI pediatric nurses and 10 caregivers were interviewed about cancer knowledge gaps, understanding of treatments, barriers to care, and preferred method of education. Based on these interviews, visual materials were identified as the most effective tools. A 20-page, 9’x12’ flipchart for use by nurses educating caregivers and patients was produced. Input from nurses and doctors led to the content used for a pilot study with 10 adult, English-speaking caregivers of UCI patients. Revisions were made to the materials based on caregiver feedback from the pilot study, and a final flipchart was printed.

Summary of results Caregivers engaged most with topics related to their family’s experience with cancer. These experiences were the basis of each topic covered in the final flipchart. After receiving one-on-one health education using the flipchart, 9 caregivers identified at least 1 sign of cancer, 8 identified at least 1 healthy food for children on chemotherapy, and 10 identified at least 1 side effect of chemotherapy. The project was presented at a meeting of 30 UCI staff, which started a conversation about expanding the project and carrying it forward, with acknowledgment that all UCI staff must be involved in educating patients.

Conclusions This project increased understanding of cancer in caregivers at the UCI and encouraged UCI staff to work together to address gaps in health literacy. Multiple departments will be involved in successfully carrying the project forward, including Pediatrics, Social Work, Emergency, and IT. Projecting the flipchart on waiting room TVs, and displaying a poster summarizing the flipchart are methods that will ensure health education is regularly delivered. Encouraging cancer survivors and their families to share their personal experiences in their villages will help spread this project’s messages in Uganda.

HELPING BABIES BREATHE – AN EVALUATION OF PROGRAM EFFECTIVENESS IN RURAL GUJARAT, INDIA

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10.1136/jim-2019-WMRC.413

Purpose of study Helping Babies Breathe (HBB) is a protocol designed for health workers to successfully navigate ‘The Golden Minute’ and reduce asphyxia-related neonatal deaths. This study aims to evaluate the effectiveness of HBB in reducing neonatal mortality in a rural community hospital in Gujarat, India.

Methods used This study took place in the Mota Fofalia Pediatric Center in Gujarat, India between April 2014 and July 2018. All birth attendants completed standardized HBB training in April 2014 and subsequent data-driven, refresher training annually. Trained observers measured HBB compliance during the study timeline using a form previously validated for rural India. Information was collected about resuscitation equipment preparation and resuscitation care provided to infants. A retrospective review of the hospital death registrar was performed to evaluate trends in stillbirth and neonatal mortality since program implementation.

Summary of results The hospital death registrar recorded a total of 2395 births from 2014 to 2018. Deliveries increased steadily from 382 to 687 respectively. A decrease in stillbirths was found after the implementation of the HBB program from 10.47 to 1.46 per thousand live births in the four-year period. An overall decline in stillbirths of 2 per 1000 each year ($R^2=0.42$) was noted. Additionally, perinatal mortality rate decreased from 13.09 to 10.19, between 2014 and 2018 respectively. Neonatal mortality rate also increased over this time period from 2.62 to 8.73 per thousand live births (table 1).

Conclusions Implementation of the HBB training program has led to a reduction in stillbirth from 2014 to 2018 in the pediatric center, lending to the idea that deaths due to previous ‘stillbirths’ may have been due to secondary apnea and not true stillbirths. Further evaluation should be pursued to evaluate the cause of the observed increased in neonatal mortality rate.

IMPACT OF PROBIOTIC ADMINISTRATION IN PRETERM INFANTS

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10.1136/jim-2019-WMRC.414

Purpose of study Poor growth in the early postnatal period is associated with long-term neurodevelopmental impairment, short stature, and metabolic disorders in preterm neonates. Probiotic Lactobacillus and Bifidobacterium strains decrease late onset sepsis, necrotizing enterocolitis, and mortality in
preterm infants, but studies of the impact of probiotics on growth have been mixed. We examined growth parameters, maximum serum alkaline phosphatase and the fecal microbiota in a cohort of preterm infants treated with either Lactobacillus reuteri 17938 (BioGaia, BioGaia AB, Stockholm, Sweden) or Bifidobacterium infantis EVC001 (Enviro, Evolve Biosystems, Davis CA).

Methods used This cohort study includes premature infants with a gestational age < 33 weeks at UC Children’s Hospital with consent obtained from the parents. All infants received either mother’s milk or donor milk. Probiotics were continued until 34 weeks. Fecal samples were collected twice weekly from infants from the time of enrollment to a corrected gestational age of 36 weeks. Change in z scores from admission to discharge was calculated for weight, length, and head circumference (HC) for each infant.

Summary of results Sequencing of the 16S ribosomal RNA gene has been completed and the bioinformatics analysis is in process for 48 very low birth weight infants. The table summarizes feeding and growth data.

Conclusions There was no difference between the two probiotic groups in growth or feeding tolerance. We are currently analyzing the fecal microbiota for differences between the two probiotic groups for α-diversity, β-diversity, and abundance of microbes associated with sepsis and necrotizing enterocolitis (e.g. Enterobacteriaceae). We are also examining correlations between growth delta z-scores and relative abundance of key bacterial taxa.

### Abstract 416 Table 1 Probiotic comparison

<table>
<thead>
<tr>
<th>L. reuteri (N=29)</th>
<th>B. infantis (N=19)</th>
<th>P value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age, weeks (median IQR)</td>
<td>28 (25, 30)</td>
<td>27 (26, 29)</td>
</tr>
<tr>
<td>First feeding, day (median IQR)</td>
<td>1 (1.2)</td>
<td>2 (1.2)</td>
</tr>
<tr>
<td>Full enteral feeding, day (median IQR)</td>
<td>14.5 (10, 24.5)</td>
<td>14 (11.5, 19)</td>
</tr>
<tr>
<td>Length of stay, days (median IQR)</td>
<td>68 (58, 111)</td>
<td>57 (46.5, 101.5)</td>
</tr>
<tr>
<td>Delta z score weight (mean SD)</td>
<td>-0.77 (0.86)</td>
<td>-0.89 (0.46)</td>
</tr>
<tr>
<td>Delta z score length (mean SD)</td>
<td>-0.82 (1.5)</td>
<td>-1.13 (1.0)</td>
</tr>
<tr>
<td>Delta z score HC (mean SD)</td>
<td>-0.85 (1.5)</td>
<td>-0.70 (1.2)</td>
</tr>
<tr>
<td>Max alkaline phos (mean SD)</td>
<td>413 (119)</td>
<td>462 (150)</td>
</tr>
</tbody>
</table>

utilized the University of Nebraska Allergen Protein Database and The Universal Protein Resource (UniProt) protein sequence database to identify a total of 2211 protein/peptide sequences.

Summary of results Each sample had between 806 and 1007 proteins/peptides, with 37 to 44 non-human proteins/sample encompassing 26 plant and animal species. Bovine proteins/peptides were the most numerous; seven unique Bos taurus proteins/peptides were found in all samples. Cat, dog, mosquito, salmon, and crab were detected in all four samples. All maternal donors ingested fish, shellfish and tree nuts. Aeroallergen proteins/peptides, including dust mite and mold were identified in all samples. Two almond proteins were detected in three samples. Two samples contained latex and chicken. One sample contained several unique proteins/peptides, including carrot, two molds (including Penicillium citrinum) and American house dust mite-like protein.

Conclusions These findings represent the first breast milk mass spectrometry analysis with identification of known allergenic proteins from food and the environment. The correlation between the exposure of allergenic protein in human milk and the development of atopic disease is unclear. This raises the question of whether breast milk can serve to induce sensitization or tolerance in infants. We are currently conducting a prospective study aiming to analyze allergenic peptides in preterm infants’ milk feeding and their risk of atopic disease later on in life.

### Abstract 415 ANALYSIS OF NON-HUMAN PROTEINS/PEPTIDES IN HUMAN BREAST MILK BY MASS SPECTROMETRY

A Hanhauser*, S Wedgwood, M Underwood. University of California Davis, Sacramento, CA

Purpose of study Human breast milk is shown to decrease the risk of atopic diseases in full term babies. But its allergenic effects on preterm babies and the etiology of its protective effect have not been established. We aim to perform a broad analysis of non-human proteins and peptides in human breast milk using mass spectrometry.

Methods used Four breast milk samples were obtained from Mommy’s Milk, a human milk research biorepository in San Diego for mass spectrometry analysis. Two samples were from preterm infant’s mothers and two were from term infants’ mothers. Mothers filled out food and environmental exposure surveys when donating their breast milk. We

Purpose of study Poor post-natal growth is associated with bronchopulmonary dysplasia and pulmonary hypertension (PH) in extremely preterm infants. In a rodent model, we have shown that postnatal growth restriction (PNGR) causes intestinal dysbiosis and PH. Human beta-defensin 2 (HBD2) is a broad spectrum antimicrobial peptide produced by many epithelial cells including enterocytes in the distal small bowel and colon. We hypothesized that HBD2 would attenuate malnutrition-associated dysbiosis and PH.

Methods used Newborn rats were assigned to a litter size of 17 (PNGR) or 10 (normal growth), and exposed to room air or 75% oxygen for 14 days. Some PNGR+hyperoxia pups were treated with recombinant HBD2 by gavage. Echocardiograms were performed on day 14 to determine the ratio of pulmonary acceleration time (PAT) to pulmonary ejection time (ET); decreased PAT:ET indicates PH. Pups were then euthanized and plasma and tissues harvested. Fulton’s index (RV weight divided by LV weight + septum weight), was quantified to identify RV hypertrophy. Lung expression of IL1β and glutathione (GSH) were analyzed as a marker of inflammation and oxidative stress respectively.

Summary of results HBD2 attenuated malnutrition-induced PH and RVH and reversed associated changes in IL1β and GSH (figure 1).

Conclusions Malnutrition causes intestinal dysbiosis leading to injury in the developing lung. Alterations in inflammatory cytokines and markers of oxidative stress are promising mechanisms. The decrease in lung IL1β with malnutrition was
unexpected and was reversed by treatment with HBD2. Investigation of changes in the gut microbiota with administration of HBD2 is warranted.

GROWTH OUTCOMES IN PRETERM INFANTS LESS THAN 1250 GRAMS AT BIRTH FED HUMAN MILK WITH HUMAN MILK BASED FORTIFIER VERSUS BOVINE MILK BASED FORTIFIER

M Chang*, L Barton, T Lin, R Ramanathan, B Cayabayb, LAC+USC Medical Center, Keck School of Medicine, University of Southern California, Los Angeles, CA; Miller Children’s and Women’s Hospital, Long Beach, CA

Purpose of study Comparison of growth outcomes in preterm infants less than 1250g at birth fed with human milk with human milk-based fortifier (EHM) versus bovine milk-based fortifier (HMF).

Methods used Retrospective data collection on preterm infants less than 1250g at birth admitted to NICU from January 2016 to May 2019 who were fed with human milk with human milk-based fortifier (EHM) (26cal/oz) or bovine milk-based fortifier (HMF) (24cal/oz). Congenital anomalies or genetic syndromes were excluded. Demographics, clinical outcomes and anthropometric measurements at birth and at discharge were collected.

Summary of results There were 53 preterm infants included in the study. EHM compared to human milk with HMF, had higher discharge weight (2925g vs 2552.5g; p=0.004), discharge weight percentiles (19% vs 4.5%; p=0.006) and discharge length (47.5cm vs 45.2cm; p=0.015). (Table 1) Extraterine growth restriction was significantly lower in preterm infants fed EHM (16% vs 60.7%, p=0.001). There was no significant difference in clinical outcomes between infants fed EHM and human milk with HMF. (Table 2)

Conclusions Preterm infants fed EHM at higher caloric intake resulted in fewer infants with extraterine growth restriction. Despite lower gestational age in infants fed EHM, the corrected gestation age at discharge was not significantly different from infants fed human milk with HMF.

GROWTH FAILURE IN NEONATES WITH GASTROSCHISIS

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Purpose of study Early growth failure is associated with later developmental delays. Growth studies in infants with gastroschisis are limited. In this multi-site study, our aim was to investigate growth failure in hospitalized infants with gastroschisis.

Methods used We included neonates with gastroschisis treated at the 5 sites in the University of California Fetal Consortium (UCFC). The UCFC developed guidelines to standardize care of infants with gastroschisis, including a feeding approach. Growth failure was characterized by changes in z-scores from birth to 14 and 30 days (d) of age, and discharge (D/C) (mild, z-score decrease of 0.8-1.2; moderate, >1.2-2; severe, ≥2; or none). Regression analysis from birth to discharge for z-scores was performed, and a mixed model was used to assess growth failure predictors.

Abstract 417 Table 1 Demographics and growth outcomes of the study population

<table>
<thead>
<tr>
<th>Outcome</th>
<th>Human milk with human milk-based fortifier (%)</th>
<th>Human milk with bovine milk-based fortifier (%)</th>
<th>p-value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestational age (median, range)</td>
<td>26 (25, 37, 7)</td>
<td>27 (26, 39, 38)</td>
<td>0.018</td>
</tr>
<tr>
<td>Birth weight (median, range)</td>
<td>2925 (1032, 3109)</td>
<td>2925 (1032, 3109)</td>
<td>0.964</td>
</tr>
<tr>
<td>Birth weight (percentile)</td>
<td>61 (56, 67)</td>
<td>61 (56, 67)</td>
<td>0.964</td>
</tr>
<tr>
<td>Length at birth (median, range)</td>
<td>49.3 (48, 50)</td>
<td>49.3 (48, 50)</td>
<td>0.964</td>
</tr>
<tr>
<td>Length at discharge (median, range)</td>
<td>54.5 (53, 58)</td>
<td>54.5 (53, 58)</td>
<td>0.964</td>
</tr>
<tr>
<td>Discharge weight (median, range)</td>
<td>54.5 (53, 58)</td>
<td>54.5 (53, 58)</td>
<td>0.964</td>
</tr>
<tr>
<td>Discharge weight (percentile)</td>
<td>15 (14, 17)</td>
<td>15 (14, 17)</td>
<td>0.964</td>
</tr>
</tbody>
</table>

Abstract 417 Figure 1 HBD2 attenuated malnutrition-induced PH and RVH and reversed associated changes in IL1β and GSH.
PATHOLOGICAL FETAL GROWTH RESTRICTION RESULTS IN REDUCTION IN LEAN MASS GROWTH THAT PERSISTS INTO THE NEONATAL PERIOD


Purpose of study Fetal growth restriction (FGR), defined by an estimated fetal weight (EFW) <10%, does not differentiate constitutionally from pathologically small fetuses. Fetuses with pathological FGR by International Society of Ultrasound in Obstetrics and Gynecology (ISUOG) criteria demonstrate reduced fat and lean mass compared to FGR fetuses who do not meet criteria. Among FGR fetuses, we hypothesized that those meeting ISUOG criteria would have reduced neonatal lean mass.

Methods used In this prospective cohort study of fetuses with an EFW of <10%, ISUOG criteria were applied to the last ultrasound prior to delivery revealing either pathological FGR (FGR-P: umbilical artery pulsatility index (PI) ≥95th%, cerebroplacental ratio (CPR) ≤5%, middle cerebral artery PI ≤5%, or EFW ≤3%; n=23) or constitutionally small (FGR-C: EFW 3-10% + normal Dopplers; n=20). Weight, height, and head circumference (OFC) were obtained at birth. Air Displacement Plethysmography (ADP) and anthropometrics with skin folds were used at 40-44 weeks to estimate lean and fat mass and were compared by linear models adjusted for postmenstrual age (PMA).

Summary of results At birth, FGR-P were born earlier (36.7 vs 38.2 wks), shorter (44.3 vs 47.4 cm), lighter (2.1 vs 2.7 kg), and had smaller OFC (31 vs 33.2 cm; P<0.01). At 40.4 wks PMA, weights were similar, but FGR-P neonates were shorter by 1.9 cm (95% CI: 0.5-3.3) and had smaller OFC by 0.8 cm (0.2-1.5) and chest circumference by 2.7 cm (0.4-5.0; P<0.05). Percent fat mass by ADP trended lower in FGR-P by 4% (P=0.08), with no difference in skin fold measurements between groups.

Conclusions FGR-P neonates had persistent reductions in length and head/chest circumference but had similar weights and fat mass compared to FGR-C neonates, indicating that catch up growth in the neonatal period in FGR-P is mostly fat mass. Increased fat relative to lean mass may contribute to future metabolic disease risk in fetuses with pathological FGR.

RACIAL/ETHNIC AND REGIONAL DIFFERENCES IN PREVALENCE AND PREDICTORS OF MEDICAL COMPLEXITY AMONG A NATIONAL COHORT OF VLBW INFANTS

KE Hannan*, SL Bourque, C Levek, S Tong, S Hwang. University of Colorado, Aurora, CO

Purpose of study To describe the prevalence and predictors (patient- and hospital-level) of medical complexity, defined as having Complex Chronic Conditions (CCC) or Technology Dependence (TD), in a nationally representative sample of very low birthweight (VLBW, <1500g) infants.

Methods used We performed a retrospective, cross-sectional analysis of hospital discharge data (2009 & 2012) from the Healthcare Cost and Utilization Project’s (HCUP) Kids’ Inpatient Database (KID). We included infants with ICD9 codes indicating birthweight <1500g and complete demographic information. Primary outcomes were CCC or TD and combined outcome of CCC or TD or death during birth hospitalization. CCC was defined using ICD9 codes indicating a medical condition expected to last >12 months and involving either several organ systems or extensive involvement of 1 organ system. TD was defined using ICD9 codes indicating medical complexity arising from dependence upon medical technology (device). All analyses were weighted to estimate national and regional prevalence of CCC, TD, and death.

Summary of results Our cohort consisted of 38,597 VLBW infants, representing a population of 78,683. A weighted percent of 39.4% (95% CI 38.9-39.8) had any CCC/TD and 56.3% (95% CI 55.9-56.7) had the combined outcome of CCC/TD or death. After adjusting for demographic characteristics, hospital characteristics, and birthweight, females (AOR 0.71, 95% CI 0.68-0.73), non-Hispanic black infants (AOR 0.92, 95% CI 0.88-0.97), and those in an urban non-teaching hospital (AOR 0.83, 95% CI 0.80-0.87) were less likely to have the combined outcome of CCC/TD or death. Compared to infants in the Northeast, those in the Midwest (AOR 1.09, 95% CI 1.02-1.15) and South (AOR 1.07, 95% CI 1.02-1.13) were more likely to experience the combined outcome, while those in the West (AOR 0.93, 95% CI 0.88-0.99) were less likely to experience the combined outcome.

Conclusions Almost 40% of VLBW infants discharged following birth hospitalization have medical complexity. Both patient and hospital-level characteristics are independently associated with CCC/TD. Understanding the prevalence and predictors of CCC/TD among VLBW infants provides critical data to inform future interventions to improve care and outcomes during hospitalization and after discharge home.

UTILITY OF BIRTH CERTIFICATE DATA FOR EVALUATING HOSPITAL VARIATION IN ADMISSIONS TO NEONATAL INTENSIVE CARE UNITS

1H Haidari, 2H Lee*, 3J Hikuzu, 4H Lin, 5X Xu, 6Stanford University, Stanford, CA; 7Yale School of Medicine, New Haven, CT; 8Yale School of Public Health, New Haven, CT

Purpose of study Efforts to study potential overuse of NICU admissions and hospital variation in practice are often hindered by lack of an appropriate data source. Birth certificates contain a data element measuring NICU admission and may offer a means for research in this area, as they...
can be linked to data sources containing diagnoses. Although multiple studies have evaluated the quality of birth certificate data elements, accuracy of the NICU admission measure remains largely unknown. To address this knowledge gap, we examined the concordance of hospital-level NICU admission rates between birth certificate data and the California Children’s Services (CCS) data to inform the utility of birth certificate data in studying hospital variation in NICU admissions.

Methods used We analyzed birth certificate data from California in 2012 and linked it to hospital-specific summary data from CCS regarding NICU admissions. NICU admission rates were calculated for both datasets while using CCS data as the gold standard. Difference between birth certificate-based and CCS-based NICU admission rates was assessed using Wilcoxon signed rank test, and concordance between the two rates was evaluated using Lin’s concordance correlation coefficient and Kendall’s W concordance coefficient.

Summary of results Among a total of 103 hospitals that were linked between the two datasets, birth certificate data generally underreported NICU admission rates compared to CCS data (7.72% vs 11.51%, p<0.001). However, in a subset of 35 hospitals where the difference in NICU admission rates between the two datasets was small, birth certificate-based NICU admission rate showed concordance with the rate from CCS data (Lin’s concordance correlation coefficient=0.91, 95% confidence interval: 0.84-0.95, Kendall’s W concordance coefficient=0.99, p<0.001). Hospitals with good concordance data did not differ from other hospitals in the institutional characteristics assessed.

Conclusions For a selected subset of hospitals, birth certificate data may offer a reasonable means to investigate hospital variation in NICU admissions.

Abstracts

Neonatology – perinatal biology II
Concurrent session
8:00 AM
Saturday, January 25, 2020

423 LATE GESTATION FETAL HYPERGLUCAGONEMIA RESULTS IN LOWER FETAL WEIGHT AND DECREASED UTEROPLACENTAL NUTRIENT UPTAKE
S Cilvik, S Wesolowski, L Brown, P Rozance*. University of Colorado Anschutz Medical Campus, Aurora, CO
10.1136/jim-2019-WMRC.423

Purpose of study Fetal glucagon concentrations are elevated in intrauterine growth restriction. Glucagon receptors are expressed in the placenta, with one previous study suggesting that glucagon may inhibit umbilical uptake of certain amino acids. We hypothesized that chronic fetal hyperglucagonemia would result in decreased amino acid transfer to the fetus and decreased growth.

Methods used Late gestation fetal sheep received a direct intravenous infusion of glucagon (5 or 50ng/kg/min; n=7 and 5, respectively) or a vehicle control (n=10). After 8-10 days, uterine and umbilical blood flows were measured utilizing the 3H2O transplacental diffusion technique. Utero-placental and umbilical (fetal) uptake rates of oxygen, glucose, lactate, and amino acids were calculated using the Fick principle. All glucagon-exposed fetuses were grouped and compared to controls using the unpaired Student’s t-test to best evaluate global trends related to hyperglucagonemia.

Summary of results Fetuses receiving glucagon had 13% lower fetal weight compared to controls (p<0.05). Fetal
MECHANISM UNDERLYING INCREASED CARDIAC EXTRACELLULAR MATRIX DEPOSITION IN PERINATALLY NICOTINE INDUCED TRANSGENERATIONAL MODEL

A Rehan*, T Chuang, J Liu, A Ansari, C Yu, R Sakurai, O Khorram, V Rehan. Lundquist Institute for Biomedical Innovation at Harbor-UCLA Medical Center, Torrance, CA

Abstracts

Purpose of study Previously, we demonstrated transgenerational transmission of perinatal nicotine-induced lung phenotype. Although perinatal nicotine exposure predisposes to offspring cardiac fibrosis, the underlying mechanisms involved and whether this phenotype is also transmitted transgenerationally are unknown. MiR-1 is known to target fibronectin and MIAT, a long non-coding RNA, is implicated in cardiac injury. We hypothesized that perinatal nicotine exposure alters the expression of miR-1 and MIAT transgenerationally.

Methods used Pregnant Sprague-Dawley rat dams received dihexit or nicotine (s.c.) 1mg/kg once daily from e6 until postnatal day (PND) 21. Pups delivered at term and breastfed ad lib. At PND21, some pup hearts were collected and some pups were weaned as breeders to generate F2 and F3 offspring, but without any subsequent exposure to nicotine. F3 hearts were collected at PND21 for analysis. For in vitro studies, neonatal primary cardiac fibroblasts (NPCF) and cardiac myocytes (CM) were isolated and treated with nicotine (10^{-9}M) for 72h to 7 days. MIAT levels were modulated by siRNA and the expression of MIAT, miR-1, and fibronectin was determined by qRT-PCR, western blotting, and immunohistochemistry.

Summary of results Perinatal nicotine exposure increased expression of fibronectin at PND21 in F1 (p<0.05; N=5) and F3 hearts (p<0.05; N=7) while decreasing miR-1 and increasing MIAT levels (p<0.05). Nicotine treatment of cultured NPCF and CM increased MIAT levels, while it suppressed miR-1 expression (p<0.05). MIAT knockdown resulted in miR-1 induction, with a decrease in fibronectin expression in cardiac fibroblasts (p<0.05).

Conclusions Perinatal nicotine exposure is associated with increased fibronectin deposition in F1 and F3 offspring, increased cardiac MIAT levels, and decreased miR-1 expression. These data provide a mechanistic basis for increased ECM deposition in the perinatally nicotine exposed heart and for the first time demonstrate the transmission of perinatal nicotine-induced predisposition to cardiac fibrosis transgenerationally.

EFFECT OF PERINATAL NICOTINE EXPOSURE ON MATERNAL/OFFSPRING GUT MICROBIOME AND SHORT CHAIN FATTY ACID PROFILES, AND THE DEVELOPING LUNG

M Martin*, 1J Liu, 1C Yu, 1Y Wang, 1R Sakurai, 1Y Liu, 1B Ji, 1V Rehan, 1Lundquist Institute at Harbor-UCLA, Torrance, CA; 2Beijing University, Beijing, China

Purpose of study Alterations in maternal gut microbiome play an important role in the development of respiratory diseases such as asthma in offspring. Although nicotine adversely affects the developing lung directly whether there is a role of altered maternal/offspring gut microbiome in mediating this effect is unknown. Since bacterial metabolites [e.g., short chain fatty acids (SCFAs)] and/or possibly the bacteria themselves cross the placental barrier, we hypothesize that the effects of perinatal nicotine exposure on the developing lung in part are mediated via alterations in maternal/offspring gut microbiome and/or their metabolic consequences. We determined the effect of perinatal nicotine exposure on maternal/offspring gut microbiome and SCFA profiles and how SCFA levels affect the key lung developmental pathways.

Methods used Pregnant Sprague-Dawley rat dams received dihexit or nicotine (s.c.) 1mg/kg once daily from e6 until postnatal day (PND) 21. Pups delivered at term and breastfed ad libitum. At PND21, dams and pups were killed to collect ceca for microbiome profiling and blood for SCFA profiling. Additionally, e14 fetal rat lung explants were cultured for 5 days ± butyric acid, propionic acid, or acetic acid (10^{-9} and 10^{-6}M for each), following which key lung developmental markers were analyzed by q-RT-PCR.

Summary of results Comparison of the relative abundance of intestinal flora among offspring at genus level showed increased lactobacillus (15 to 49%), but decreased bacteroides (22 to 9%) abundance in the nicotine exposed group (N=3; p<0.05). Consistent with these data, concentrations of fecal SCFAs, especially butyric acid, which is produced predominantly by bacteroides were reduced in both maternal and offspring sera. Fetal lung explants cultured in lower concentrations of SCFAs (10^{-9}M vs 10^{-6}M) resulted in lower expression of lung maturation markers such as PPARγ and surfactant protein B (p<0.05). Conclusions Perinatal nicotine exposure results in altered offspring gut microbiome, especially, decreased abundance of SCFAs producing bacteroides, which provides a novel mechanism underlying perinatal nicotine exposure induced alterations in lung maturation and development.
ENDOTOXEMIA IMPAIRS NEONATAL AND JUVENILE HEPATIC SELENOCYSTEINE SYNTHESIS AND SELENENOZYME DEFENSE
K Sjostrom*, L Sherlock, L Nguyen, M Zarate, E Nozik-Grayck, C Wright. University of Colorado, Aurora, CO
10.1136/jim-2019-WMRC.426

Purpose of study Pediatric sepsis is a major cause of childhood morbidity and mortality. Nutritional status contributes to host response after sepsis, and selenium (Se) is an essential micronutrient with potential to improve outcomes during life-threatening infection. Se levels decrease during sepsis and inversely correlate with risk of multiorgan dysfunction, pneumonia, and death. However, Se replacement has not improved outcomes in septic adults. The liver is primarily responsible for converting Se into the bioactive amino acid selenocysteine (Sec). We have previously demonstrated endotoxemia decreases hepatic factors for Sec synthesis in adult mice, which may obstruct the therapeutic potential of Se replacement during illness. It is unknown if this phenomenon is experienced in younger mice. We hypothesized endotoxemia would impair hepatic Se processing both in neonatal and juvenile mice.

Methods used Male and female P7 and P21 C57/B6 mice were exposed to IP LPS (5 mg/kg) and sacrificed at 0, 6, and 24 hr. Hepatic mRNA and protein expression for factors essential in Se processing (PSTK, SepsecS/SLA, Sephs2/Scly and EEFsec, Scly) and selenoenzymes (SelenoP, Gpx1, Msrb1, Trxr1) were measured by qPCR and Western blot.

Summary of results LPS decreased hepatic mRNA of factors for Sec synthesis at 6 hr, including PSTK, SepsecS, Sephs2, Scly and EEFsec, in P7 and P21 mice (p<0.05, n=4-5). At 24 hr, mRNA remained low only in the P7 mice (p<0.05, n=4-5). Protein expression for Sephs2 decreased at 24 hr in P7 and P21 mice. Protein for PSTK decreased at 6 hr at both ages and remained low at 24 hr only in the P7 mice (p<0.05, n=4-5). Hepatic transcription of SelenoP, Gpx1, Msrb1 and Trxr1 decreased at 6 and 24 hr in P7 mice. SelenoP and Msrb1 mRNA decreased at 24 hr in P21 mice (p<0.05, n=4-5).

Conclusions Endotoxemia downregulates numerous hepatic factors for Sec synthesis and selenoenzymes in P7 and P21 mice. This downregulation demonstrates delayed recovery in P7 mice. We speculate that neonates may be more susceptible to hepatic oxidative stress during infections due to an exaggerated decrease in hepatic Sec and selenoenzyme production.


UMBILICAL CORD MILKING WITH VENTILATION IN PERINATAL ASPHYXIA AND MECONIUM ASPIRATION
M Harde*, A Lesneski, P Vail, S Lakshminrusimha. UC Davis, Davis, CA
10.1136/jim-2019-WMRC.427

Purpose of study Placental transfusion is currently not recommended for neonates in need of resuscitation at birth due to concerns about delaying positive pressure ventilation (PPV). Umbilical cord milking (UCM) during PPV with an intact cord may potentially combine the benefits of placental transfusion without delaying the onset of PPV. The objective of this study was to evaluate pulmonary and systemic hemodynamics and gas exchange following UCM and PPV in a model of birth asphyxia, meconium aspiration syndrome (MAS) and persistent pulmonary hypertension of the newborn (PPHN).

Methods used Twelve near-term gestation (142/150d) lambs were asphyxiated by umbilical cord occlusion. MAS was induced by instilling meconium into the lungs during gasping leading to PPHN. Lambs were randomized to UCM+PPV or immediate cord clamping (ICC). Lambs were ventilated for an hour and preductal blood gases, left pulmonary (Qlpa) and carotid (Qca) blood flows were monitored.

Summary of results Baseline characteristics were similar between the groups. Following ICC, Hgb decreased after birth. With UCM+PPV, Hgb was stable and O2 delivery to the brain was significantly higher compared to the ICC group. Qca decreased and Qlpa increased with ventilation in both groups.

Abstract 427 Table 1 Comparison of ICC followed by PPV vs. UCM+PPV before cord clamping—mean (SEM)

<table>
<thead>
<tr>
<th>Group</th>
<th>Immediate cord clamping followed by PPV (n=6)</th>
<th>Umbilical cord milking + PPV before cord clamping (n=8)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Birth weight (g)</td>
<td>3065 (422)</td>
<td>3210 (495)</td>
</tr>
<tr>
<td>Gender – Male (%)</td>
<td>2 (33%)</td>
<td>3 (50%)</td>
</tr>
<tr>
<td>Multiplicity (single/twins/duplet)</td>
<td>2/3</td>
<td>2/4</td>
</tr>
<tr>
<td>Fetal asphyxia gas pH</td>
<td>6.92 (0.03)</td>
<td>6.92 (0.07)</td>
</tr>
<tr>
<td>Fetal asphyxia lactate (mmol/L)</td>
<td>6.7 (0.7)</td>
<td>7.1 (0.4)</td>
</tr>
<tr>
<td>Baseline Hgb (g/dL)</td>
<td>13 (3.8)</td>
<td>11.6 (0.8)</td>
</tr>
<tr>
<td>Baseline fetal carotid flow (mL/kg/min)</td>
<td>28 (4)</td>
<td>33 (5)</td>
</tr>
<tr>
<td>Baseline fetal pulmonary flow (mL/kg/min)</td>
<td>32 (12)</td>
<td>34 (7)</td>
</tr>
<tr>
<td>PaO2/PaCO2 at 5 min after delivery</td>
<td>203 (26)</td>
<td>205 (44)</td>
</tr>
<tr>
<td>Carotid flow (mL/kg/min)</td>
<td>15 (2.1)</td>
<td>20 (2.9)</td>
</tr>
<tr>
<td>Pulmonary flow (mL/kg/min)</td>
<td>91 (24)</td>
<td>93 (13)</td>
</tr>
<tr>
<td>Body Temperature (°C)</td>
<td>36.3 (0.3)</td>
<td>36.8 (0.4)</td>
</tr>
<tr>
<td>Blood Gas pH</td>
<td>7.23 (0.04)</td>
<td>7.19 (0.05)</td>
</tr>
<tr>
<td>Blood Gas Hgb (g/dL)</td>
<td>13.9 (1.6)</td>
<td>12.3 (0.6)</td>
</tr>
<tr>
<td>Blood Gas Lactate (mmol/L)</td>
<td>3.4 (0.7)</td>
<td>3.6 (0.4)</td>
</tr>
<tr>
<td>Arterial oxygen content (CaO2) (mL/dL)</td>
<td>14.1 (4.0)</td>
<td>15.1 (0.7)</td>
</tr>
<tr>
<td>Oxygen delivery to the brain (mL/kg/min)</td>
<td>2.1 (0.2)</td>
<td>3.0 (0.3)*</td>
</tr>
<tr>
<td>Oxygen extraction by the brain (%)</td>
<td>25.2 (2.9)</td>
<td>17.6 (2.4)</td>
</tr>
</tbody>
</table>

Conclusions Lambs with perinatal asphyxia tolerated UCM +PPV with an intact cord did not have a decrease in Hgb and had higher oxygen delivery to the brain at 60 min after birth compared to lambs with immediate cord clamping.

CONTINUOUS CHEST COMPRESSIONS WITH ASYNCHRONOUS VENTILATIONS INCREASE CEREBRAL BLOOD FLOW
10.1136/jim-2019-WMRC.428

Purpose of study Current guidelines recommend interrupted chest compressions (CC) at a 3:1 compression-to-ventilation (CV) ratio in severely bradycardic newborns. In neonates, heart rate is the primary determinant of cardiac output and achieving a higher rate of CC may increase blood flow. We
hypothesize that continuous CC with asynchronous ventilations (CCCaV) leads to quicker return of spontaneous circulation (ROSC) and better hemodynamics compared to 3:1 CV resuscitation.

Methods used Sixteen near-term lambs (142/147d) were asphyxiated by cord occlusion to cardiac arrest. Lambs were randomized to 3:1 CV or CCCaV (120 CC/min and asynchronous ventilations at 30/min). After 5min of asystole ventilation was provided by a T-piece resuscitator with CC started after 30s. First dose of epi was given at 6min if ROSC was not achieved.

Summary of results There were no differences in baseline characteristics between groups (table 1). Incidence and time of ROSC was similar in both groups (table 1). Lambs that received CCCaV had significantly greater left carotid blood flow compared to 3:1 CV (8.1 ±3.3 vs 5.3 ±2.5 ml/kg/min, p=0.01; figure 1). There was no difference in systolic, diastolic and mean BP between groups.

Purpose of study Trophoblast stem cells (TSCs), the precursors to placental differentiated cells, have unique self-renewal properties and have the ability to differentiate into many trophoblast cell subtypes. These qualities make TSCs important to study as they are highly involved in development and placenta functioning. Recent studies have identified certain genes that are involved in the differentiation of TSCs. Through overexpression of these genes, it may be possible to reprogram differentiated cells back into their TSC state.

Methods used To study the possible effects of these various genes, plasmid constructs were made with our genes of interest. RNA from cell lines rich in TSCs was used to make a cDNA library, from which the cDNAs for the target genes were amplified by PCR. These amplified genes were cloned into a retroviral expression E. coli shuttle vector using the Gibson Assembly method. Midscale plasmid purification was performed and the DNA was transduced into fibroblast cells to validate the function of the constructs.

Summary of results Our initial qPCR results indicated that these transduced genes were well expressed in the fibroblast and HEK293 cells. With confirmed expression of these genes, we need to determine if there is also production of protein products, and if these transcription factors lead to reprogramming of the fibroblast cells into induced TSCs (iTSCs).

Conclusions Upon successful reprogramming, iTSCs can be used to study gene expression patterns in the differentiation of TSCs into their differentiated daughter cells. This will also allow for greater understanding of placental-related pregnancy complications.

Abstract 428 Table 1

<table>
<thead>
<tr>
<th>Group</th>
<th>3 to 1 (n=8)</th>
<th>CCCaV (n=8)</th>
<th>p value</th>
</tr>
</thead>
<tbody>
<tr>
<td>Weight kg (SD)</td>
<td>3.9 (1.1)</td>
<td>4.0 (0.9)</td>
<td>0.90</td>
</tr>
<tr>
<td>Sex M:F</td>
<td>5:3</td>
<td>4:4</td>
<td>1</td>
</tr>
<tr>
<td>Time to asystole (min)</td>
<td>15.1 (5.4)</td>
<td>13.1 (2.4)</td>
<td>0.37</td>
</tr>
<tr>
<td>ROSC</td>
<td>7/8 (88%)</td>
<td>7/8 (88%)</td>
<td>1</td>
</tr>
<tr>
<td>Time to ROSC</td>
<td>5.1 (2.4)</td>
<td>6.1 (1.5)</td>
<td>0.32</td>
</tr>
<tr>
<td>ROSC without Epi</td>
<td>3/8 (38%)</td>
<td>3/8 (38%)</td>
<td>1</td>
</tr>
</tbody>
</table>

Abstract 428 Figure 1

Conclusions In this model, CCCaV showed greater carotid blood flow compared to 3:1 CV resuscitation. Clinical studies assessing the neurodevelopmental outcomes comparing CCCaV to 3:1 CV during newborn resuscitation are warranted.

Purpose of study Various sulfonamides are protective in lung ischemia-reperfusion injury both by carbonic anhydrase inhibition and other mechanisms.

Methods used 30 healthy male rats (300-350 g, 6-8 weeks old) were randomly classified into six groups: 1. Sham (n=5). 2. I/R injury (n=5) treated with saline. 3. I/R injury + 30
mg/kg BZ pretreatment (n=5). 4. I/R injury + 30 mg/kg AZ pretreatment (n=5) and 5. I/R injury + 30 mg/kg NMA pretreatment (n=5). Rats in the sham group underwent left thoracotomy without any hiliar clamp or treatment. For the IR injury groups, the left main pulmonary artery and bronchus were clamped for 60 minutes and then blood flow was reestablished for 90 minutes, after which measurements were taken and tissue samples obtained.

Summary of results Az, NMA, but not BZ, significantly reduced the fall in PaO$_2$/FiO$_2$, a measure of gas exchange efficiency, caused by lung I/R injury. The rise in lung wet to dry weights (reflecting pulmonary edema), and protein extravasation (reflecting capillary permeability) were reduced by all pretreatments. The lungs in pretreatment groups showed less neutrophilic infiltration, alveolar edema and hemorrhage. Lastly, drug pretreatments all reduced HIF-1 activation, suggestive of less tissue hypoxia with I/R injury.

Conclusions Our data demonstrate that pretreatment with AZ, BZ and NMA all protect against and mitigate the extent of damage with I/R in the lung. The protection appears to involve direct effects of CA inhibition, but in the case of NMA, other actions independent of CA inhibition that may include either anti-oxidative or hypoxia-mediated Ca$^{++}$signalling changes, as has been shown in the uninjured hypoxic pulmonary vasculature.

### Abstracts

#### 431 INTENSIVE REGISTERED RESPIRATORY THERAPIST ATTENTION MAY REDUCE LENGTH OF STAY AND READMISSION RATE IN ACUTE EXACERBATIONS OF COPD

N Sarma*, M Dolan, K Craddock, S Louie, J Kuhn, J Schivo. UC Davis Health, Sacramento, CA; VA Medical Center, Sacramento, CA

Purpose of study Acute exacerbations of chronic obstructive pulmonary disease (AECOPD) are common and lead to significant morbidity and mortality. Institutional AECOPD quality measures such as hospital length of stay (LOS) and readmission rates are increasingly important. Prior studies demonstrated benefit from patient education (e.g. inhaler technique), appropriate medications (e.g. mist inhalers for patients with reduced inspiratory ability), and a follow up plan. Bundling these proven methods, our institution implemented a Registered Respiratory Therapist (RRT)-directed program, identifying AECOPD patients and personalizing management. We aim to analyze the effect of this RRT program on hospital LOS and readmissions as well as the improvement in patient symptoms over time.

Methods used We will perform a retrospective case control study comparing a well-described cohort of AECOPD patients who received intensive RRT attention to AECOPD patients who received standard care. Intensive RRT attention is defined as daily inpatient visits by an RRT COPD Case Manager, inculcating patient education, customized medication selection, and a follow up plan. Usual care includes a one-time RRT evaluation with medication recommendations. We aim to analyze the 800 subjects per group enrolled over the last 6 years to show a reduction in LOS by 0.5 days in the intensive group. The power analysis of our cohort size is acceptable to detect a difference of 10% using a paired t test. We will also compare readmission rates and COPD Assessment Tool (CAT) scores.

Summary of results We expect a reduction in LOS by 1.5 days and readmissions among AECOPD patients in the intensive RRT program. We also expect an improvement (decrease) in symptoms for patients who receive appropriate medications and instructions.

Conclusions Use of a dedicated COPD RRT program will likely significantly improve meaningful clinical and institutional quality outcomes in AECOPD patients.

#### 432 PROVISION OF SMOKING CESSION RESOURCES IN THE CONTEXT OF LUNG CANCER SCREENING

1) Shen*, 1,2)M Triplette. 1University of Washington School of Medicine, Seattle, WA; 2Fred Hutchinson Cancer Research Center, Seattle, WA

Purpose of study Evidence-based lung cancer screening (LCS) with low-dose chest CT is now widely recommended for high-risk current and former smokers. The potential benefits of screening may go beyond early cancer detection, as shared-decision making (SDM) presents a ‘teachable moment’ to reinforce cessation and provide resources for current smokers. However, there are few studies on smoking cessation practices in the context of LCS in a clinical setting. The objective of this study was to understand smoking cessation initiation and practices during LCS SDM encounters.

Methods used This is a retrospective study of participants in a multi-center LCS program with a SDM clinical encounter who were current smokers at time of screening (n=469). The LCS tracking database and EHR were manually reviewed for patient demographics and clinical encounter information. The Charlson Comorbidity Index (CCI) was determined through automated review of all ICD-10 codes. The primary outcome was provision of smoking cessation resources, defined as: documented referral to cessation resources, referral for nicotine replacement and/or provision of prescription medication. Participant and provider factors associated with the primary outcome were evaluated using chi-squared testing and multivariable logistic regression.

Summary of results The majority of study participants were men (70%), with a median age of 62 years and smoking history of 50 pack-years. The median CCI was between 2 and 3. 71% of encounters had some discussion of smoking, but only 21% of participants were referred to tobacco service, 31% were provided nicotine replacement resources and 17% were provided related prescriptions. In a multivariable model, the provision of smoking cessation resources was half as likely in participants with higher levels of comorbidity CCI>2 (OR: 0.58, 95% CI: 0.37-0.90).

Conclusions Overall provision of smoking cessation resources among current smokers was low, particularly among patients with higher levels of comorbidity. Interventions are needed to improve smoking cessation counseling and resource utilization at the time of LCS encounter, and this may be especially important in patients and providers with chronic illness who face competing priorities for time and resources.
Methods used The UC Davis Environmental Health Sciences Center launched an online survey covering a region affected by the 2018 Camp Fire and the lingering heavy smoke. The survey launched in May 2019, seven months post-fire. We then attempted to recruit survey respondents from a list of 137 respondents within Butte County, most with persistent respiratory symptoms. Those we reached were asked to attend a one-day pilot research clinic in Butte County on June 15, 2019. On short notice, the research team was assembled with three clinical research coordinators, two respiratory therapists, a UC Davis faculty physician, and an internal medicine resident. The protocol included validated respiratory symptom questionnaires, spirometry, measurement of exhaled nitric oxide, and exhaled breath condensate collected with our novel handheld device and stored for subsequent analysis.

Summary of results 22 patients were scheduled, 20 of whom attended and consented to undergo clinical examination. 8 participants had an Asthma Control Questionnaire score >10, and 6 had an Asthma Control Test score <20, suggesting significant respiratory symptoms in the past month. Among those subjects, only 1 had a forced expiratory volume in 1 second (FEV1) <80% and an FEV1/FVC (forced vital capacity) of <70%. Exhaled nitric oxide concentrations were elevated >30 ppb in 2 subjects, which corresponds to a specificity for asthma diagnosis of 0.84 and a sensitivity of 0.53.

Conclusions This clinic demonstrated a strong response with a low no-show rate and feasibility of the protocol. The data suggest that we can use this approach to identify both incident asthma and longitudinal respiratory symptom burden following wildfire exposure.

435 DERIVATION AND VALIDATION OF A DIAGNOSTIC PREDICTION MODEL FOR INTERSTITIAL LUNG DISEASE

Objective Derivation of a prediction model for idiopathic interstitial (ARDS) using electronic health records.

Methods A prediction model was derived and validated using electronic health records from the University of California, Davis, and a community hospital.

Results The model included variables such as age, smoking status, and pulmonary function tests, with a receiver operating characteristic area under the curve of 0.85.

Conclusions The model is a promising tool for predicting ARDS.
ILD. Variables used to construct the ILD risk score included age, PFT indications, height, total lung capacity, forced expiratory volume in 1 second and diffusion capacity. Similar test performance was observed across cohorts, with the ILD-Screen having a sensitivity of 0.82 and specificity of 0.8 when applied prospectively. ILD-Screen positivity increased the odds of ILD by 19-fold (OR 18.9, 95% CI 13.6-26.4) and explained a significantly higher proportion of ILD risk (0.81) when compared to common ILD features, including cough (0.58), dyspnea (0.53) and lung crackles (0.66).

Conclusions The ILD-Screen has good sensitivity and specificity for detecting ILD. Systematic implementation of the ILD-Screen has potential to reduce diagnostic delays and facilitate early intervention. The benefit of such approach must be weighed against the cost and risk of false positives, which will be common due to the low prevalence of ILD. Further research is needed to determine the clinical implications of its use.

Case reports II
Concurrent session
10:15 AM
Saturday, January 25, 2020

436 UNUSUAL PRESENTATION OF ACUTE HEPATITIS COMPPLICATED BY PANCREATITIS IN A CHILD
OC Dogan*, M Haight. UCSF Fresno, Fresno, CA
10.1136/jim-2019-WMRC.436

Case report A 2 year-old previously healthy boy presented with 1 day of nonbilious nonbloody emesis and diarrhea, in the setting of cough, congestion, and rhinorrhea for 1 week. Parent gave a total of Acetaminophen 1600 mg over 1 week, and 1 tablet of Aspirin 4 days prior to the presentation. Examination showed small ecchymosis on sternum. Initial labs included complete blood count, complete metabolic panel, lipase, toxicology screen, coagulation studies. They showed elevated values for white blood count at 17000/µL, alanine transaminase 2467 U/L, aspartate aminotransferase 3985 U/L, lipase 3302 U/L, amylase 1167 U/L, international normalized ratio 1.3, and prothrombin time 16.6 seconds. Salicylate, acetaminophen, and bilirubin levels were unremarkable. Investigations for infectious causes showed EBV DNA PCR of 18604 copies/ml with negative hepatitis A, B, and C, viral respiratory panel and stool pathogens. Abdominal computerized tomography scan showed mild pericholecystic fluid. Abdominal ultrasound was unremarkable. Hereditary pancreatitis panel, CMV PCR and HSV PCR were negative. He was subsequently diagnosed with acute hepatitis complicated by pancreatitis without cholestasis secondary to EBV infection.

This is an unprecedented presentation of acute hepatitis complicated by pancreatitis secondary to EBV infection in a child. In infants and young children, clinical course of EBV infection is generally silent. When complicated by hepatitis, EBV infection does not cause cholestasis, hyperbilirubinemia, or jaundice in younger populations. Our patient had viral prodromal symptoms of cough, congestion, and rhinorrhea for 1 week, and then presented with emesis and diarrhea. Laboratory results revealed highly elevated levels of white blood count, transaminases, lipase, and amylase, and was diagnosed with acute hepatitis with pancreatitis. Infection with EBV may cause a viral prodromal episode with a complication of hepatitis. EBV-induced hepatitis with pancreatitis is very unusual and has been reported in 4 patients in the literature between ages of 8 to 11 years. This case is significant in that it illustrates the importance of investigations of infectious agents in children presenting with acute hepatitis with pancreatitis, as it influences further management and outcome prediction.
MOYAMOYA IN PREGNANCY, RETROSPECTIVE CASE REPORT AND LITERATURE REVIEW

BA Mocan*, F Venter, M Gill, N Idemudia, S Jung, D Bell, C D’Assumpcao, S Ragland, K Sabetian. Kero Medical-UCLA, Bakersfield, CA

10.1136/jim-2019-WMRC.438

Purpose of study Moyamoya disease is a rare chronic occlusive cerebrovascular disease characterized by progressive stenosis at the terminal portion of the internal carotid artery and an abnormal vascular network at the base of the brain. Clinical manifestations include transient ischemic attacks, cerebral infarcts, and intracranial hemorrhages. Pregnancy invokes intravascular volume and intracerebral pressure changes from conception to delivery. Here we discuss the management of a pregnant patient diagnosed with moyamoya disease after stroke four years prior who presented to us with a transient ischemic attack.

Methods used Retrospective case report.

Summary of results A 24 year old G1P0 at 34 weeks gestation presented with sudden inability to swallow, speak, and right finger tingling lasting 10 minutes. She has residual left arm and facial weakness from a stroke four years prior and was diagnosed with Moyamoya disease at that time. Magnetic resonance imaging found no acute infarction or hemorrhage but showed old right frontotemporal encephalomalacia and old left caudate lacunar infarct. Magnetic resonance angiography found attenuation and irregularity of distal right and left internal carotid arteries with nonvisualization of the right distal internal carotid artery at the level of the circle of Willis and abrupt severe narrowing of the left internal carotid artery near the circle of Willis. There was severe attenuation of the right and left anterior and middle cerebral arteries. 16 hour continuous EEG was normal with no interictal epileptic discharges or seizure activity. There was concern for her proceeding with labor and vaginal delivery specifically in regards to valsalva causing increased intracranial pressure, which could cause intracranial hemorrhage, and hyperventilation, which could lead to vasoconstriction of small collateral blood vessels and cause cerebral infarct. Therefore the patient had cesarean section with no intra- or post-operative complications.

Conclusions Moyamoya disease in pregnancy requires awareness of physiological changes during labor and delivery, making cesarean section a favorable choice for optimal hemodynamic maintenance. Multidisciplinary team is recommended.

CRONOBACTER SAKAZAKII BACTEREMIA AND MENINGITIS IN A 4-WEEK-OLD FEMALE

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10.1136/jim-2019-WMRC.439

Background Cronobacter sakazakii is a foodborne pathogen that can cause life-threatening bloodstream infections and meningitis predominantly in neonates. It is associated with CNS complications, including abscess and cyst formation. We present a case of a neonate infected with this bacterium.

Case This is a 4-week-old female born at full term who presented with 1 day of fever, fussiness, and poor oral intake. The patient had 1 week of nasal congestion and had sick contacts at home. No vomiting, diarrhea, or rashes were reported. Birth history revealed that the mother was GBS positive during pregnancy but was treated appropriately before delivery. On exam, the patient was febrile, ill-appearing, irritable, and tachycardic.

A full sepsis workup was performed. Ceftriaxone was started empirically. Initial labs revealed a WBC of 19,000, sodium of 133 and CSF which showed a bloody tap with RBC of 250,000/mmc, WBC of 250/mmc, 79% polymorphonucleocytes, protein of 279mg/dL, glucose of 64mg/dL. Urinalysis, Influenza/RSV swab, and CXR were unremarkable. Blood and CSF cultures grew Gram-negative bacilli that were speciated as Cronobacter sakazakii. Although the organism was sensitive to ceftriaxone, the antibiotic was switched to cefepime for concerns of resistance; gentamicin was added for a synergistic effect. MRI of the brain showed an arachnoid cyst, and serial imaging did not show further CNS complications. A repeat lumbar puncture proved CSF sterility and gentamicin was discontinued. Subsequent blood cultures were negative for growth. Fevers, irritability, oral intake, WBC, and electrolytes improved after starting antibiotics. The patient completed a 21-day course of IV antibiotics and was discharged home.

Discussion Cronobacter sakazakii infections are rare. The bacterium is highly virulent with a mechanism to survive even in stressful environments such as very dry climates. The CDC reports that the mortality rate for Cronobacter meningitis may be as high as 40%. It has been linked to the ingestion of contaminated powdered infant formula. Formula samples were sent to the Health Department for testing which returned negative. Cronobacter sakazakii should be considered as a pathogen for neonates with meningitis who live in arid environments in addition to the commonly expected organisms.

SPONTANEOUS CEREBELLAR ECTOPIA IN A PATIENT WITH COCCIDIOIDAL MENINGITIS

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10.1136/jim-2019-WMRC.440

Introduction Spontaneous cerebellar ectopia without acute trauma is rare. Meningitis has not yet been reported as a cause in the literature. We present a case of coccidoidal meningitis complicated by cerebellar ectopia.

Case report 22 year old Hispanic female with five year history of coccidoidal meningitis complicated by non-communicating hydrocephalus requiring ventricular peritoneal shunt with multiple revisions who is on posaconazole and biweekly intrathecal amphotericin B deoxycolate through an ommaya reservoir presented with acute encephalopathy. Two days prior to presentation, she had lumbar puncture for prognostic studies. She developed new onset headache, confusion and then somnolence, for which family brought patient to hospital. Imaging suggested shunt malfunction due to worsening hydrocephalus as the distal VP shunt was looped around the bladder. She developed fever upon admission. VP shunt and ommaya were removed and an external ventricular device was placed. Samples from VP shunt grew Enterococcus faecalis. The following day she became obtunded requiring intubation. Neuroimaging found obstructive hydrocephalus with compression on each side of the brainstem and downward pressure on the cerebellar tonsils, which extend 13mm below the foramen magnum. She exhibited right sided decorticate and left sided decerebrate posturing as well as right medial gaze palsy. EEG was...
consistent with coma. EVD drainage was decreased from 20mmHg to 5mmHg and patient was placed in head-down position. She did not show any signs of improvement and 8 days later new VP shunt was placed. The following day she began to track with eyes and follow simple commands. One week later tracheostomy was placed and due to VP shunt malfunction underwent another revision. PEG was placed. After 23 ICU days, patient recovered enough to be stable for telemetry floor. After 6 days, she was transferred to acute rehab. At four months follow up, patient has no neurological deficits, performing all ADLs and is able to drive a vehicle.

Conclusions Cerebellar ectopia is not common and when it occurs the sequelae is permanent and prognosis is very poor. Spontaneous appearance and resolution without neurological sequelae is rarely seen and the mechanism is unknown.

441 A RUPTURED MYCOTIC ANEURYSM IN A 17-YEAR-OLD MALE WITH STAPHYLOCOCCUS LUGDUNENSIS ENDOCARDITIS

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Introduction Staphylococcus lugdunensis is coagulase-negative staphylococcus (CoNS). Like other CoNS, it is considered a normal skin flora but can cause life threatening infections, including endocarditis. Mycotic aneurysm is a rare complication of endocarditis with reported mortality up to 80%. There have been limited case reports of mycotic aneurysms with Staphylococcus lugdunensis endocarditis. We report the first pediatric case of Staphylococcus lugdunensis endocarditis complicated by a mycotic aneurysm in a 17-year-old male.

Case report A 17-year-old male presented with a one month history of cough, fever, night sweats, weight loss, and 2 weeks of right leg pain with difficulty ambulating. His exam was notable for a Grade 2/6 systolic murmur. Doppler study was unremarkable for a deep venous thrombosis of the right lower extremity. Blood culture resulted positive for Staphylococcus lugdunensis. Echocardiogram demonstrated moderate mitral regurgitation with mitral valve prolapse concerning for endocarditis. He was initially treated with Vancomycin then deescalated to IV Cefazolin based on susceptibilities. On day 12 of illness, he developed stabbing pain in his right anterior medial thigh and was found to have a tender, pulsatile mass. Doppler study revealed a femoral artery mycotic aneurysm. CT angiogram showed occlusion of the right profunda femoris artery with ruptured pseudoaneurysm. He underwent emergent embolization of the right profunda femoris artery without complications. Clinically he did well and completed a 6-week course of IV Cefazolin. Due to mitral valve prolapse with regurgitation he is a candidate for future valve replacement.

Discussion Staphylococcus lugdunensis associated mycotic aneurysm is a rare but serious complication requiring immediate intervention. To date, there have been two reported cases with Staphylococcus lugdunensis endocarditis complicated by mycotic aneurysm with one resulting in death due to rupture. Early diagnosis and intervention is critical in improving outcomes.

Conclusion This case illustrates the rare presentation of a mycotic aneurysm secondary to Staphylococcus lugdunensis endocarditis and reinforces the importance of early diagnosis and intervention to improve outcomes.
Abstracts

444 MEDIC! A COURSE ON IMPROVISED MANAGEMENT OF TRAUMA PATIENTS FOR MEDICAL STUDENTS
Z Smith*, University of Washington, Seattle, WA
10.1136/jim-2019-WMRC.444

Purpose of study Physicians may be the first responders in a trauma situation when away from their clinic and equipment, and anecdotal reports prove this to be a real possibility. Considering the variable response time of emergency medical services, there is a need for training on trauma management in a low resource, out-of-hospital setting. Through their education medical students potentially have an advantage over the lay bystander, if provided adequate instruction. Searches via PubMed articles, civilian emergency medical services, military combat medicine guidelines and the author’s own Special Forces Medic training. Instruction stressed hands-on skills practice with both classmates and mannequins as patients.

Methods used An electronic survey was distributed to current osteopathic medical students from all years of training across the nation. The survey was comprised of 14 questions in total, with four questions pertaining to responder demographics, six questions designed to assess knowledge in concussion diagnosis and treatment, and four questions assessing exposure to previous concussion education. Data was collected in aggregate. Results by year of training in medical school were analyzed via one-way ANOVA. Multiple comparison combinations between all four years of training were assessed via Tukey’s HSD test.

Summary of results Preliminary collection of over 400 responses and analysis of the data show a statistically significant linear correlation between year of training in medical school and success in answering questions regarding correct identification of concussion symptoms, proper diagnosis, and management protocols. Interestingly, a large majority of respondents indicated that they wanted to receive more formal concussion education, with a small minority feeling that they have received adequate concussion education to prepare them for their clinical years as a physician.

Conclusions While data collection is still ongoing, the preliminary results of our study suggest that as medical students progress in their years of training, so too does their knowledge of concussion diagnosis and management. However, both the total performance results on the knowledge assessment, and the results of the students’ self-assessment of concussion knowledge, demonstrate that students at all academic levels lack a comprehensive understanding of concussion symptoms and management. The data from our study suggests both a desire and need from medical students to strengthen concussion education in medical schools across the country.

Summary of results Success was measured through instructor observation of techniques during class and student feedback following the trial’s conclusion. Evaluations revealed significant increase of an average 4 points on a 10-point scale in self-reported confidence levels performing various skills. Students’ comments revealed that they overwhelmingly valued the course content’s applicability and the amount of practical and hands-on training. The most consistent criticism was the desire for a longer class period (90 instead of 60 minutes).

Conclusions Creation of this course was part of a new curriculum development option for satisfying a medical school research requirement. The course content fulfills a gap in medical training, and the trial sessions of Medic! demonstrated this course’s viability among medical students. It reinforces core curriculum lessons, prepares them to respond appropriately in a trauma situation, and instills confidence in the no-equipment, out-of-hospital setting. Future plans for the course are to conduct a full iteration during the author’s fourth year of medical school.

445 OCHO PASOS: A QUALITY IMPROVEMENT PILOT TO IMPROVE BREASTFEEDING SUPPORT IN OUTPATIENT CLINICS
P Mendosa*, T Hatfield, J Martinez, D Sedillo, T Ozechowski, E Sebesta, MM Ramos. University of New Mexico; Albuquerque, NM
10.1136/jim-2019-WMRC.445

Purpose of study The documented health benefits of breastfeeding (BF) for mother and baby are numerous, yet a third of US mothers achieve only three months of exclusive BF despite the American Academy of Pediatrics (AAP) recommendation to exclusively BF for six months. National experts have recently published evidence-based guidelines for outpatient clinics to support longer durations of BF. Our pilot study sought to evaluate the feasibility of an intervention to improve support for BF in outpatient primary care clinics.

Methods used We compared evidence-based BF-friendly guidelines from the AAP, Academy of Breastfeeding Medicine, California’s 9 Steps, and Washington’s 10 Steps to create Ocho Pasos (Eight Steps) to tailor these guidelines to the New Mexico population. In 2018, we recruited an outpatient primary care clinic in rural New Mexico to participate in a 6-month pilot program using the Ocho Pasos as a framework to evaluate the clinic based on provider, staff, and patient interviews. After establishing a baseline, we focused on improving medical record documentation of BF supportive care through two Plan-Do-Study-Act (PDSA) cycles. Approximately 40 medical records were reviewed at baseline and at each subsequent medical record review (MRR).

Summary of results Providers and staff set a goal to improve documentation of mothers’ intended BF duration. Documentation improved from 0% at baseline to 63.0% and 91.3% at the second and third MRR, respectively. A majority (16/17, 94.1%) of a convenience sample of mothers surveyed after their infants’ well child checks reported discussing their intended duration of BF with their provider. Overall, providers and staff reported the quality improvement initiative was feasible and acceptable in terms of processes and instruments; however, staff did request that fewer medical records be reviewed to reduce the time burden.
Conclusions Our preliminary findings suggest that quality improvement initiatives aimed to implement evidence-based practices for BF support in an outpatient setting are feasible. We plan to refine and expand this initiative to other outpatient clinics in New Mexico.

**446** THE IMPACT OF TELEMEDICINE ON PSYCHIATRY CONSULTATIONS

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10.1136/jim-2019-WMRC.446

**Purpose of study** Synchronous telepsychiatry (STP) and asynchronous telepsychiatry (ATP) are two modes of telemedicine used in clinical practice to improve patient access to specialty psychiatric care. However, little is known about the types of clinical recommendations given by the psychiatrists in these visits and the response of the primary care physician (PCP) to those recommendations. This study aims to characterize and compare these recommendations in ATP versus STP.

**Methods used** This study is a secondary analysis of a randomized clinical trial comparing the effectiveness of ATP versus STP. Using the Consultation Liaison Outcome Evaluation System as a foundation, expert physicians and telemedicine researchers created an expanded framework for classifying psychiatrist recommendations and rating PCP adherence to those recommendations. Recommendations were classified into five groups (‘Medication,’ ‘Diagnosis Action,’ ‘Psychotherapeutic,’ ‘Follow-up Timing,’ and ‘Other’) and thirteen subgroups. After extracting recommendations from the baseline psychiatrist visit note, blinded reviewers rated PCP adherence to those recommendations (‘Fully’, ‘Partially’ or ‘Not’) using electronic medical record notes in the subsequent six months.

**Summary of results** To date, preliminary data have been abstracted on the baseline visits of 122 patients: 62 in ATP and 60 in STP. The patients were on average 34 years old (SD=15.4) and most were already taking a psychiatric medication (84%). Psychiatrists made an average of 6.4 recommendations in the baseline consultation (range, 2-14). In comparison to STP visits, ATP visits had a greater median number of recommendations (84%), meaning higher adherence to those recommendations. There were no statistically significant differences in the discussion of the five recommendation groups and the 13 subgroups. For example, medication recommendations were made in all the ATP visits (100%) and most of the STP visits (97%).

**Conclusions** Preliminary results indicate that there may be slight differences between the amount of recommendations in ATP and STP; and little variation in the recommendation profiles. Compared to similar literature, this study found a similarly large emphasis on medication recommendations. Future work will focus on comparing the adherence of the PCPs to the recommendations between STP and ATP.

**448** RISK STRATIFICATION IN HOSPITALIZED CHILDREN WITH MEDICAL COMPLEXITIES (CMC) & HEALTH OUTCOMES

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10.1136/jim-2019-WMRC.448

**Purpose of study** At a national level, half of pregnant women receive the flu vaccine and 54% receive the Tdap (Tetanus, Diphtheria, and Pertussis) vaccine as part of their prenatal care. We assessed vaccine literacy and uptake by pregnant women to identify potential knowledge gaps.

**Methods used** In summer 2019, we surveyed the generally low-income, English-speaking patients at UCLA-Harbor Medical Center Obstetrics clinic in Torrance as part of an IRB-approved (LABioMed and Western University) exempt protocol. Patients age ≥ 18 years at ≥ 32 weeks gestational age were approached. After verbal consent was obtained, the survey of 31 questions was administered on a one-to-one basis for 6 weeks.

**Summary of results** A total of 97 patients was approached; 92 participated. 51% of subjects were age 20-27 (age range 18-50); 52% were Hispanic/Latino, 25% African American, 5% Caucasian. Overall, 75% reported receiving the Tdap vaccine; 65% said they would accept the flu vaccine. When asked if they understood why the Tdap vaccine was important during pregnancy, 22% chose ‘No’ and 14% chose ‘I Don’t Know’. The primary reason (70%) for no vaccination was not remembering having been offered Tdap. Other explanations included religious conflicts and safety concerns. Of the 35% who would not receive the flu vaccine, 44% said they believed it is ‘not effective,’ 78% said they were concerned with getting sick or having side effects, and 69% said they were concerned about safety risks it posed to themselves.

**Conclusions** This population had higher vaccination rates than national published levels (75% vs 54% Tdap and 65% vs 50% Flu); however, there continues to be a substantial gap, especially for the flu vaccine. We found patients reported a deficiency in physician discussion regarding the Tdap, suggesting a need for improvement in vaccine education. Moreover, patients may have misconceptions regarding the flu vaccine that play a larger influence on their decisional process. To improve vaccine uptake, different educational strategies should be tailored to each vaccine.

**447** PRENATAL VACCINE KNOWLEDGE AND UPTAKE AMONG PREGNANT PATIENTS

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10.1136/jim-2019-WMRC.447

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admission (TA), Length of Stay (LOS), number of Readmission (RR) (@ 3 & 6 months) prior to the study period were collected.

A statistical sample independent t-test analysis compared risk scores & utilization outcomes.

Summary of results CMC -Medical Device-19 of 40 (47.5%) CMC

19.5% of patients with high risk SDH.

Conclusions Complexity index & special needs scoring is important in risk stratification as it has a direct predictive effect on outcomes.

CMC patients have 3.5 times greater complexity & 1.5 times higher degree of disability. The results demonstrate increase health utilization with 3 times longer LOS & twice the (RR) over 180 days than the non-CMC patient group.

The longer LOS even when compared to national data from a 2006 National study is high, 7.5 vs 5.2 days. Surmissable contributing factors maybe related to higher social determinants (19.5%), poor access to care (22% of patients without PCP), & high technology dependence (47% for CMC).

**Hematology and oncology III**

Concurrent session

10:15 AM

Saturday, January 25, 2020

**Abstract 448**

| Table 1 Risk based tools: degree of complexity & effect on outcomes |
|---------------------------------|---------------------------------|
| SCORES (Mean results) | Children with Complex (n=40) | Non Complex Children (random) (n=40) |
| Exeter H.O.M.E.S. (highest complexity & utilization) | 7.55 out of 10 | 2.10 out of 10 |
| (Most complex and high utilization) | Standard Deviation 2.04 | Standard Deviation 2.18 |
| Special Needs Score (high disability) | 3.63 out of 5 | 2.0 out of 5 |
| Standard Deviation = 1.15 | Standard Deviation = 1.31 |
| Number of Readmission (in 6 months) after discharge | 2.1 standard deviation = 1.77 | 1.3 Standard deviation = 0.47 |
| Number of Readmission (90 days) after discharge | 1.0 Standard deviation = 1.29 | 1.4 Standard deviation = 0.59 |
| Length of Stay (LOS) | 7.5 days National Average (2006) = 5.2 days* | 2.1 days |

Conclusions The novel PCGC at SCCA was established to provide prostate cancer patients with genetic counseling, germline testing, combined with comprehensive discussion about potential treatment options and family implications. It facilitated immediate and future treatment options for men with prostate cancer, provided consultative services and access to critical information for patients and families about clinical trials.

Hematology and Oncology III

Concurrent session

10:15 AM

Saturday, January 25, 2020

**449**

**PROSTATE CANCER GENETICS CLINIC: GERMLINE GENETIC TESTING IN PRACTICE AND ITS IMPLICATIONS FOR PRECISION THERAPY AND FAMILY COUNSELING**

D Pov*, A Sokolova, H Cheng, University of Washington, Seattle, WA

**Purpose of study**

Prostate cancer has a significant heritable component, notably in genes involved in homologous recombination DNA repair: *BRCA2*, *BRCA1*, and others. The importance of identifying men who carry these inherited mutations is two-fold: for treatment decision making and for family counseling. The Prostate Cancer Genetics Clinic (PCGC) was established in September 2016 at Seattle Cancer Care Alliance (SCCA) to provide prostate cancer patients with cancer risk assessment, genetic testing, recommendations for treatments and clinical trials. Here, we describe the development of this unique clinic, patient demographics, and results of genetic testing for patients seen during the first 21 months since establishment. The aim of this study is to broadcast our experiences to the broader oncology community with the goal of fostering the development of similar clinics.

**Methods used**

A protocol was designed and approved by the institutional review board to retrospectively collect and analyze basic demographic data and clinical information of patients seen at PCGC between September 2016 and June 2019.

**Summary of results**

A total of 160 patients were seen between September 2016 and June 2019. 142 patients had germline testing performed, 44/160 (25%) patients had a pathogenic or likely pathogenic variant in DNA repair genes involving homologous recombination (*BRCA2*, *BRCA1*, *ATM*, and *CHEK2*). All 44 patients were advised to pursue family testing and all 31 patients with variants in homologous recombination genes were counseled on treatment options. At the time of last follow up (data abstraction period was July – August 2019), 3/31 (10%) were started on platinum-based chemotherapy or a PARP inhibitor as a direct consequence to germline testing, and 12/44 (27%) with known germline mutations confirmed at least one family member had undergone cascade genetic testing.

**Conclusions**

The novel PCGC at SCCA was established to provide prostate cancer patients with genetic counseling, germline testing, combined with comprehensive discussion about potential treatment options and family implications. It facilitated immediate and future treatment options for men with prostate cancer, provided consultative services and access to critical information for patients and families about clinical trials.
CAUSES OF HOSPITALIZATION AMONG CANCER PATIENTS RECEIVING IMMUNE CHECKPOINT INHIBITORS

A Choi*, G In, K Brady. LAC+USC Medical Center, Los Angeles, CA

10.1136/jim-2019-WMRC.450

Purpose of study As increasing number of patients receive immune checkpoint inhibitors (ICI) as cancer therapy, there is growing recognition of the immune related adverse events (IRAE) which may be serious and life-threatening. Here we report our institution’s retrospective analysis of patients treated with ICI who were hospitalized due to IRAE.

Methods used Patients with advanced cancers and who received ICI therapy at the Los Angeles County Medical Center were identified from a pharmacy database. Patients who were hospitalized within 1 year from their last dose of ICI therapy were included for analysis. Patients admitted for cancer-related complications or scheduled procedures were excluded. Toxicities were graded by CTCAE V5. Suspected immune related adverse events (sIRAE) were defined as hospitalizations where IRAE was: 1) the primary diagnosis, 2) included as a differential diagnosis, 3) biopsy proven, or 4) improved with immunosuppressive therapy.

Summary of results Among a total of 192 cancer patients receiving ICI, there were a total of 107 patients and 213 hospitalizations chosen for analysis. Lung cancer was the most common malignancy treated (25.2%). Among these 107 patients, 54.0% were on ICI at time of hospitalization, and 86.5% of hospitalizations occurred within first 6 months after initiating ICI. Among all hospitalizations, 15% required ICU level care while 10.3% required PCU level care.

Among all 213 hospitalizations, 12.2% were due to sIRAE. Among this subset of patients with sIRAE, 19% underwent biopsy, 54% received steroids, and 4% received other immunosuppressive therapy. Melanoma was the most common cancer associated with sIRAE, followed by gastrointestinal, lung, genitourinary and gynecologic cancers. Among patients with sIRAE, 69.2% were on ICI at time of admission, and 11.5% required ICU level care. The most common organ systems involved by sIRAE included: gastrointestinal (including liver), pulmonary, hematologic, endocrine, and rheumatologic.

Conclusions IrAE can affect any organ system and are likely under-diagnosed due to unfamiliarity with novel ICI therapy. IRAE can be serious and may lead to hospitalization, and may also result in life-threatening toxicity requiring ICU level care. IrAE should be on the differential for all patients hospitalized during or even after the use of ICI.

EMERGENCY-RELEASE BLOOD TRANSFUSIONS AFTER POSTPARTUM HEMORRHAGE AT THE INTERMOUNTAIN HEALTHCARE HOSPITALS

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10.1136/jim-2019-WMRC.451

Purpose of study Hemorrhage is the leading cause of child-birth-associated maternal death worldwide and is a prominent cause of morbidity and mortality in the United States. Most women do not have crossmatched blood immediately available to treat massive postpartum hemorrhage. Therefore, an emergency-release blood transfusion (ERBT) service is critical for all obstetric patients. We conducted a quality improvement project to assess our emergency postpartum transfusion practices, as a basis for progress.

Methods used We collected data retrospectively from January 2011 to December 2018 at all Intermountain Healthcare hospitals, on ERBT after postpartum hemorrhages; logging circumstances, number and type of transfusion product, and outcomes.

Summary of results The cohort comprised 224,035 live births. Ninety women received ERBT for postpartum hemorrhage (4.02 transfused women/10,000 live births). The four most common conditions listed as casually associated with ERBT were uterine atony (39%), abruption/previa (16%), retained placenta (11%), and uterine rupture (7%). The mean number of component units transfused per woman was 7.8. About 75% of the transfused women (67/90) received ≥3 units of blood products. The Trauma Service recommend ratio of blood products administered for massive hemorrhage is either 1:1:1 or 2:1:1 (RBC:FFP:platelets), however our ratios varied widely. Only 1/67 women receiving ≥3 units had a 1:1:1 ratio, and only 6/67 received a 2:1:1 ratio, thus the remaining 90% (60/67) received an inadequate dose of FFP and platelets, compared to that recommended for acute massive hemorrhage. Nineteen% (17/90) of women had a hysterectomy, 34% (31/90) had an ICU admission, and 1 died.

Conclusions Emergency transfusions for postpartum hemorrhage occurs in 1/2500 births. Thus, obstetric-specific ERBT policies are imperative for all hospitals with obstetrical services. Transfusion of RBC alone is insufficient to provide hemostasis during massive ongoing hemorrhage. For postpartum hemorrhage, we encourage further study of the resuscitative and hemostatic benefits of a balanced ratio of blood products or transfusing low-titer group O cold-stored whole blood.

CHANGES IN BLOOD COUNT PARAMETERS MAY OFFER EARLY INDICATORS OF CLONAL HEMATOPOIESIS OF INDETERMINATE POTENTIAL (CHIP)

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10.1136/jim-2019-WMRC.452

Purpose of study CHIP is associated with increased risk for developing hematologic neoplasms and cardiovascular disease. Genetic mutations underlying CHIP include epigenetic modifiers such as DNMT3A, TET2, and ASXL1 (also known as DTA mutations), which are also found in myelodysplastic syndromes (MDS) and acute myeloid leukemia (AML). DTA mutations are also associated with lower likelihood of relapse when detected in AML patients in remission.

Inflammation may be a unifying factor between clonal hematopoiesis and cardiovascular disease, as hematopoietic stem cells proliferate during systemic inflammation, and atherosclerosis involves chronic inflammation. Because inflammatory marker elevations are strongly associated with red blood cell distribution width (RDW), we explored whether RDW and mean corpuscular volume (MCV), which are both often...
ABSTRACTS

Elevated in MDS, may predict hematologic neoplasm progression. Methods used Records of 45 patients seen for pre-leukemic conditions who underwent next gen sequencing were analyzed for CHIP-associated gene mutations, particularly DTA mutations. MCV and RDW were also compared between patients with and without DTA and other mutations. Summary of results Those patients who had ≥1 DTA mutation (31/45; 69%) had more all CHIP mutations (mean: 3.29, SD: 1.35) than those with no DTA mutations (mean: 1.43, SD: 0.65). ASXL1, DNTM3A, SRSF2, TET2 and TP53 were most frequent (≥22% for each). MCV differed between DTA (mean: 96.1, median: 93.9, range: 62.5-117.8, SD:11.3) and non-DTA patients (mean: 91.4, median: 92, range: 80.8-102.2, SD: 7.5), as did RDW (DTA: mean: 19.8, median 19.2, range: 12.2-36.5, SD: 5.1; non-DTA: mean 16.5, median 16.3, range: 13.2-20.6, SD: 2.1). Conclusions DTA mutations are associated with lower likelihood of AML relapse, f AML relapse, but are only found through expensive genetic sequencing. Our early results suggest that differences in MCV and RDW may help predict patients more likely to have DTA mutations. However, the presence of confounding conditions causing microcytosis, including iron deficiency, would mitigate the trend toward elevated MCV in patients with DTA mutations. Thus, routine CBCs may be an inexpensive indicator of patients who would benefit from genetic testing to better characterize disease and stratify risk of developing more aggressive hematologic disorders.

Purpose of study Cutaneous squamous cell carcinoma (CSCC), the second most common type of skin malignancy, occurs in patients with chronic sun exposure or who are immunocompromised. Though PD-1 blockade is FDA approved for the treatment of advanced and metastatic CSCC, much remains to be elucidated regarding the clinical implications of CSCC and the toxicities that may manifest from treatment. Methods used We report one institution’s experience treating CSCC with PD-1 blockade. We conducted a retrospective chart review among patients who received PD-1 blockade for CSCC. Tumor responses were evaluated by RECIST 1.1. Summary of results Among 18 CSCC patients receiving PD-1 blockade, median age was 62 years (range 22-92), and 13 (72.2%) were male. Six patients (33.3%) had tumors on the head and neck, 5 (27.8%) on the lower extremity, 4 (22.2%) in the anorectal region, 2 (12.5%) on the torso, and 1 (6.25%) on the upper extremity. Median tumor mutational burden was 10.5 mutations per Mb (range 5-177). Four (22.2%) patients had positive PD-1 expression, using a 1% cutoff. Seventeen patients were evaluated by RECIST. Three patients (17.6%) had a complete response, 3 (17.6%) had partial response, and 2 (11.7%) had stable disease, yielding an overall response rate of 35.3% and a disease control rate of 47.1%. Nine (52.9%) patients had progression of disease. Of the patients with a clinical response, the median number of cycles of PD-1 blockade received was 4 and the median number of cycles to response was 2. Median time to response was 10.6 weeks, median progression free survival (PFS) was 21.6 weeks, and median duration of response was 30.3 weeks. Nine (50%) of patients had toxicities of any grade, with 3 patients (16.7%) having Grade 3 or higher toxicity. Five patients (17.6%) died during our study period, 4 from disease and 1 from toxicity. Conclusions PD-1 blockade induces responses in approximately 35% of patients with advanced CSCC, a finding consistent with major clinical trials demonstrating its efficacy. While PD-1 blockade was generally well tolerated, some patients did develop serious, life-threatening toxicity.
Conclusions Comparable fingerprints of NHL subtypes using a micro-Raman system and probe provide support for potential diagnostic use of the RESpect probe in a clinical setting and encourages future use of RESpect in the setting of pediatric cancers.

**Abstracts**

**Abstract 455 A CASE OF BILATERAL RENAL MASS IN BECKWITH-WIEDEMANN SYNDROME: NEPHROMEGALY VS NEPHROBLASTOMATOSIS**

J Lim*, R Natarajan, Z Mehta, H Cheam. University of Las Vegas School of Medicine, Las Vegas, NV

10.1136/jim-2019-WMRC.455

Summary A baby boy who was born at 36 weeks and 5 days, weighing 4160 grams via emergency cesarean section due to decreased fetal cardiac activity and a biophysical profile of 2/10 to a G1P1 mother was found to have bilateral renal masses, cardiac hypertrophy, and polyhydramnios on prenatal ultrasound. Upon delivery, he was found to be limp, unresponsive, and apneic. Positive-pressure ventilation was initially started but required intubation at 1 minute of life. Umbilical vein catheter was also placed around 2.5 minutes of life. A 30 mL normal saline bolus was given with improvement in tone, color, and respiratory effort. Initial blood gas was concerning for HIE. He was noted to have features that were concerning for Beckwith-Wiedemann syndrome (BWS) which included macrosomia, macroglossia, nephromegaly, and hypoglycemia. Abdominal MRI revealed diffuse enlargement of both kidneys with replacement of the parenchyma by numerous homogeneous confluent masses, compatible with nephroblastomatosis. CT of the abdomen with contrast revealed bilateral nephromegaly, but no discrete masses to confirm nephroblastomatosis was seen. Upon further discussion with a COG nephrologist, they believed it was nephromegaly associated with BWS.

Discussion Nephromegaly has been shown to be a high risk factor in developing nephroblastomatosis and Wilms tumor in patients with BWS. Nephroblastomatosis is the presence of diffuse nephrogenic rests, which are clusters of embryonic metanephric cells. Its finding is significant, as it is considered to be a precursor of Wilms tumor. It has been found in 100% of kidneys with bilateral Wilms tumor and 40% with unilateral Wilms tumor. Although, over 90% of the patients with BWS will not develop a tumor, the presence of nephromegaly warrants close monitoring. The only associated risk factor for developing a tumor is in patients with hemihypertrophy. It has been suggested that even with no evidence of nephroblastomatosis on ultrasound, it can be missed if they have microscopic nephrogenic rest which may later develop into Wilms tumor. With the possibility of missing nephroblastomatosis or different interpretations of imaging, close follow-up with interval imaging should be considered in patients with nephromegaly in the setting of BWS.

**Immunology and rheumatology II**

**Concurrent session**

**10:15 AM**

**Saturday, January 25, 2020**

**Abstract 456 RISK OF ELIMINATION DIETS IN ATOPIC DERMATITIS PATIENTS WITH FOOD HYPERSENSITIVITY**

1M Sun*, 2T Sierro, 3A Armstrong. 1University of California, San Diego, La Jolla, CA; 2University of Southern California, Los Angeles, CA

10.1136/jim-2019-WMRC.456

Purpose of study While studies have examined the utility of diet therapy in unselected atopic dermatitis (AD) populations, there is a critical gap of knowledge regarding the potential risks of elimination diets in patients with AD and food hypersensitivity. Therefore, we sought to examine the risks of elimination diets in AD patients with food hypersensitivity.

Methods used We performed a systematic review of primary literature based on patients with AD and food hypersensitivity. We extracted studies from PubMed, EMBASE, and CENTRAL from inception to July 2019. We assessed the quality of these studies using a risk of bias tool appropriate for each study type. Randomized control trials were assessed with the Cochrane Risk of Bias Tool and observational studies were assessed with the Newcastle-Ottawa scale.

Summary of results A total of nine papers with 1227 participants with atopic dermatitis was analyzed. The quality of observational studies ranged from fair to poor and that of...
interventional studies from low to high risk of bias. The domains assessed for bias in the interventional studies appear in table 1. The most concerning risks associated with elimination diets are anaphylaxis and growth restriction.

Conclusions There are clear risks associated with elimination diets; however, the range of severity and prevalence of these risks have not been clearly established. Patients with AD should be properly diagnosed with food allergens before adopting elimination diets. Additionally, AD patients should be counseled on the risks associated with elimination diets regardless of their food hypersensitivity status. Ultimately, whether or not the risks associated with elimination diets outweigh the benefits in AD patients with food hypersensitivity cannot be generalized and must be determined on an individual basis.

### Abstracts

#### 457 OMEPRAZOLE INDUCED SUBACUTE CUTANEOUS LUPUS ERYTHEMATOSUS

**Purpose of study** Drug induced subacute cutaneous lupus (SCLE) is an uncommon condition and a form of Drug-induced lupus erythematosus and it is well described in the literature. Many drugs can trigger the condition including proton pump inhibitors. There are several physical finding clues to the diagnosis. We are describing a severe form of SCLE associated with high dose omeprazole.

**Methods used** Retrospective Study

**Summary of results** Here we present a case of a 34-year-old Hispanic female with a persistent refractory acid re flux disease, erosive esophagitis and gastritis. She underwent fundoplication 5 months prior and about 2 weeks later she developed dysphagia, odynophagia, circum-lip lesions, arthralgia and myalgia. In about 3 weeks prior she developed skin lesions starting on her back and spread throughout her body. The lesions were pruritic and started to ulcerate with sub-ungal hematomas. Her painful oral lesions worsened and white material supper impos. Antibiotics and antifungal did not relieve and she came and was admitted. Her medication history was significant for high dose omeprazole of 40 mg twice day through out as the only medication she was taking. Her physical exam showed severe buccal mucositis superimposed with candida and ulceration on her lips similar to steven johnson syndrome. She had diffuse erythematous base vesicular and pustular lesions in various stages. She also had sub-ungual hematomas to 4th digit of the left hand and hallux of left foot with periungual telangiectasia. Her laboratory studies showed His-tone Ab 1.6 U (1.6-2.5) and rest of her work up for rheumatological and infectious disease came back negative including SS-A and SS-B. Diagnosis of drug induced SCLE were made and she was started on steroid with resolution of her symptoms to be followed up in clinics on taper dose of steroid. Her omeprazole were discontinued.

**Conclusion** Drug induced subacute cutaneous lupus should be considered in the appropriate clinical setting the main remedy is to avoid insulting agent and use of corticosteroids.

#### 458 NEONATAL NET-INHIBITORY FACTOR INHIBITS MACROPHAGE EXTRACELLULAR TRAP FORMATION

**Purpose of study** Extracellular traps (ETs) are web-like structures composed of chromatin and anti-microbial peptides extruded from immune cells to trap and kill microbes. Neutrophil extracellular traps (NETs) participate in antimicrobial defense; however, dysregulated NET formation leads to inflammatory tissue damage in many disease states including sepsis. Neonatal NET-Inhibitory Factor (nNIF) is a novel peptide found in fetal circulation that inhibits NET formation. Macrophages also form extracellular traps (METs), and while less well described, METs are pathogenically implicated in numerous disease states and may contribute to inflammatory tissue damage. Whether nNIF inhibits extracellular traps from cells other than neutrophils remains unknown. We hypothesize that nNIF inhibits MET formation.

**Methods used** We isolated murine alveolar macrophages via bronchial alveolar lavage. Isolated macrophages were stimulated with lipopolysaccharide (LPS; 100 ng/mL) to induce MET formation ± nNIF pre-incubation (10 nM) for 1 hour in vitro. MET formation was qualitatively assessed using live cell imaging with cell permeable (SYTO green) and cell impermeable (SYTOX orange) DNA stains. We also performed this experiment in surrogate macrophages differentiated from an immortalized human monocyte cell line - THP-1 cells (PMA 20 nM for 2 days).

**Summary of results** LPS induced in vitro MET formation in both murine alveolar macrophages and surrogate human macrophages as assessed qualitatively and quantitatively. Furthermore, preincubation with nNIF reduced in vitro MET formation in both cell types.

**Conclusions** We conclude that nNIF inhibits MET formation in vitro by murine and surrogate human macrophages. We speculate that inhibition of MET formation by nNIF may lead to a reduction in inflammatory tissue damage in a variety of disease states, including sepsis.

#### 459 CD20 EXPRESSION IS ELEVATED ON A T HELPER CELL SUBPOPULATION IN NEWLY DIAGNOSED MULTIPLE SCLEROSIS PATIENTS

**Purpose of study** Anti-CD20 monoclonal antibody therapy (rituximab, ocrelizumab) has demonstrated significant efficacy in treating relapsing remitting multiple sclerosis. Although it is postulated that these therapies’ mechanism of action is to deplete peripheral B cells—the primary antigen presenting cells to effector T cells in the MS disease process—we sought to understand if CD4+/CD40+ T helper (Th40) cells, which are elevated in autoimmune states including T1D and MS may also be targeted by the drug.

**Methods used** Frozen peripheral blood mononuclear cell (PBMC) samples from 10 newly diagnosed MS patients were cultured in vitro with one of either IL-2, IL-3, or Interferon-γ, and then stimulated by either biotinylated CD3
or biotinylated CD3 and CD28. We stained these samples with anti-CD3, -CD4, -CD40, and -CD20 antibodies, then conducted flow cytometry at baseline and 24 hours post-stimulation.

**Summary of results** CD20 expression on Th40 cells was significantly elevated compared to the broader population of coexpressive CD40+CD4+ T Cells both at baseline and 24 hours (Mean diff. = 22.9%, 95% CI: 20.4% to 25.3%, p<0.0001; Mean diff. = 17.5%, 95% CI: 15.8 to 19.2%, p<0.01, respectively). ANOVA demonstrated CD20 expression was also significantly elevated on all CD4+ cells compared to CD3+ cells (F(1, 408) = 126.3, p<0.01 at baseline and 24 hours (Mean diff. = 6.2%, 95% CI: 5.1% to 7.2%; Mean diff. = 6.9%, 95% CI: 6.0% to 7.7%, respectively). CD20 expression decreased on CD4+ cells between baseline and 24 hours (Mean diff. = -2.2%, 95% CI: -3.3% to -1.1%, p<0.01) and on CD3+ cells between baseline and 24 hours (Mean diff. = -2.9%, 95% CI: -3.7% to -2.1%).

**Conclusions** This research demonstrates that CD20 is preferentially expressed on cells that coexpress CD4 and CD40 (Th40) cells in MS patients. Because Th40 cells are present at elevated proportions in patients newly diagnosed with multiple sclerosis and are implicated in autoimmune disease progression, these findings may indicate an alternative mechanism of action for anti-CD20 monoclonal antibody therapy. Next steps will include analyzing CD20+ Th40 cell counts in patients before and after commencing anti-CD20 antibody therapy for multiple sclerosis.

**Summary of results** Compared to PAN implant CFH, age-matched controls displayed significantly reduced rod response amplitudes. Analysis of tissue sections showed an accumulation of complement C3 in the retina of age-matched controls, with retinal thinning, and photoreceptor atrophy.

**Conclusions** Here we present a novel intravitreal PAN implant capable of slowing disease progression in an AMD animal model. This research is critical in determining a method of halting AMD, ultimately preventing blindness in individuals who suffer from the disease. In addition to AMD, the PAN implant may potentially be used to treat other retinal degenerations mediated by inflammation and CF, including retinal dystrophies and diabetic retinopathy. The results of this study will be critical in guiding pre-clinical studies of the PAN implant and ultimately the first human clinical trial of the PAN implant.
Purpose of study Coccidioidomycosis is a fungal infection that is endemic to California, Arizona, and other areas of the southwestern United States. It is caused by inhalation of spores of Coccidioides immitis. 60% of infected patients are asymptomatic; the remaining 40% have primarily pulmonary disease characterized by fever, cough, and pleuritic chest pain. In rare cases, less than 1% of infected patients, dissemination of the infection can occur. Dissemination usually affects those with impaired cellular immunity and pregnant women, and can involve bones, joints, meninges, and/or skin [1].

We present a very rare case of disseminated polyarticular coccidioidomycosis that presented with three isolated joints in an otherwise healthy 29-year-old male.

Methods used Retrospective case report.

Summary of results We present the case of a 29-year-old male who presented to the emergency department (ED) complaining of pain and swelling in multiple joints for three months. In the ED, an arthrocentesis of two of the joints involved were significant for total nucleated cells of 520,000/cm² and 93% Neutrophils respectively. Serum anti-Coccidioides IgM and IgG titers were greater than 1:512. Whole body bone scan showed bony involvement of the affected joints. Based on the history obtained and lab results collected, patient was diagnosed with a rare case of disseminated polyarticular Coccidioidomycosis and osteomyelitis. Subsequently he was treated with long term Amphotericin B, after which, he showed clinical improvement.

Conclusions This case demonstrates an atypical presentation of disseminated Coccidioidomycosis. It illustrates the challenges in approaching a presentation of multiple possible etiologies while highlighting the necessity of considering Coccidioidomycosis in the differential diagnosis of a patient presenting with polyarthritis in endemic areas. Furthermore, it shows that arthrocentesis with an extremely elevated WBC count and neutrophil predominance, as opposed to the expected lympho-dendritic predominance, should raise suspicion of Coccidioidomycosis in these endemic areas.

Neonatology general VI
Concurrent session
10:15 AM
Saturday, January 25, 2020

CONGENITAL DIAPHRAGMATIC HERNA: IMPROVEMENT IN SURVIVAL WITHOUT ECMO FOLLOWING PRACTICE CHANGE
M Yang*, CC Yost, S Fenton, B Yoder. University of Utah/Primary Children’s Hospital, Salt Lake City, UT

Conclusions CDH guideline changes focused on minimizing stimulation, pre-ductal oxygenation, and less aggressive ventilator/inotrope support were associated with decreased ECMO use and improved survival without ECMO.
TELEMEDICINE CONSULTATIONS MAY IMPROVE STABILITY OF NEONATES PRIOR TO TRANSPORT


10.1136/jim-2019-WMRC.464

Purpose of study Telemedicine can improve quality of care for neonates born in rural community hospitals. However, no studies to date have examined the effect of telemedicine on neonatal stabilization prior to and during the transfer process. The purpose of this study was to explore the association between neonatal telemedicine consultations and infant stability during transfer, compared to traditional telephone consultations.

Methods used We collected data from all telemedicine and telephone consultations for newborns that took place at six community hospitals in northern California between 2014 and 2018. To measure infant stability, we used a modified Transport Risk Index of Physiologic Stability version II (TRIPS-II) from the California Perinatal Transport System (CpTS), a validated instrument designed to measure illness severity in newborns. Scores were collected at three different time points for infants who were transferred: at the time of consultation, time of arrival of the transport team, and time of arrival at the regional neonatal intensive care unit (NICU). We performed difference-in-difference analyses to examine changes in TRIPS-II scores between different time points for infants receiving either telemedicine or telephone consultations.

Summary of results A total of 209 infants were transferred to a higher level of care after receiving either type of consultation. 162 infants (77.5%) received a telemedicine consult and 47 (22.5%) received a telemedicine consultation. At the time of consultation, infants who received telemedicine had higher mean TRIPS-II scores than those who received a telephone consultation (10.9 vs 5.5, p=0.01). Infants who received telemedicine consultations experienced a mean decrease in TRIPS-II scores of 3.31 points between time of consultation and time of transport team arrival, compared to a decrease of 0.2 points for infants who received a telephone consultation only (p=0.007). There was no difference in TRIPS-II scores upon arrival to the NICU.

Conclusions Our findings suggest that telemedicine may help to stabilize neonates at local hospitals prior to the arrival of transport. This may have important implications for the use of telemedicine for neonates requiring transfer.

STABILITY OF DARBEPOETIN IN NEONATAL INTRAVENOUS FLUIDS

1U Nwokidu-Aderibigbe*, 2L Jantzie, 1J Maxwell, 3S Beauman, 4RK Ohls. 1University of New Mexico, Albuquerque, NM; 2Johns Hopkins University, Baltimore, MD; 3University of Utah, Salt Lake, UT

10.1136/jim-2019-WMRC.465

Purpose of study We previously reported stability of erythropoietin in commonly used neonatal intravenous (IV) fluids. Since that time, Darbepeotin (Darbe) has been introduced as a long-acting erythropoiesis stimulating agent. We determined the stability of Darbe over time in common neonatal IV fluids and determined the effect of the excipient stabilizing agent Polysorbate 80 (PS80) on Darbe recovered.

Methods used The following solutions were studied:

- 0.9% normal saline (NS), dextrose 10% in water (D10), dextrose 10% in water + 4.5% amino acids (DA), and standard total parenteral nutrition (TPN). PS80 was added to each solution to a final concentration of 0.05% (w/v). Darbe was diluted with each solution (with and without PS80 added) to a goal concentration of 50 pg/mL. Each solution was drawn into a 10 mL syringe and pushed through 60-inch neonatal microbore tubing. Samples were collected at baseline (prior to push), 5 minutes, 1 hour and 24 hours (for TPN only), and frozen immediately at -80°C. Darbe concentrations were subsequently measured in duplicate via the Quantikine ELISA Human Erythropoietin Immunoassay.

Summary of results Darbe concentrations were 32.7±1.4% of expected when diluted with NS, D10 or DA solutions without PS80. When PS80 was added to maintain the original excipient concentration, baseline concentrations were 96.2±1.8% of their expected (p<0.01 versus non-PS80 solutions). Concentrations remained similar to baseline at the 5 and 60-minute time points. Conclusions A significant decrease in the expected concentration of Darbe occurred in fluids without PS80. Darbe was most stable in TPN over time, despite a lower than expected baseline concentration. We speculate that adding to TPN solution once weekly might be a novel and sufficient way to administer Darbe to neonates intravenously.

IN-ROOM TECHNOLOGY EXPEDITES TREATMENT OF NEONATAL SEIZURES IN THE INTENSIVE CARE NURSERY


10.1136/jim-2019-WMRC.466

Purpose of study Seizure burden in the neonatal period is independently associated with poor neurodevelopmental outcomes. Given the potential for seizure-induced brain injury in the neonate, seizures should be treated promptly. We aimed to decrease the time to medical treatment of neonatal seizures from 25 minutes (pre-intervention) to less than 15 minutes.

Methods used The study is set in a 58-bed level IV Intensive Care Nursery (ICN) at an academic children’s hospital. Previously, a Seizure Rescue Task Force including a team of physicians, nurses, pharmacists and technicians was created as a hospital-wide initiative to expedite seizure treatment in hospitalized pediatric patients. The seizure rescue process was implemented in Dec 2016 and included a seizure rescue button added to the in-room staff terminal to alert a pharmacist for bedside expedited administration of anti-seizure medication. Hospital-wide impact was significant but the ICN had poor adoption. We created a key driver diagram to identify unique barriers to the process in the ICN. In Jan 2019 we rolled out an education initiative about the importance of rapid treatment of neonatal seizures and use of the seizure rescue process. We reviewed the medical records of all patients in the ICN with seizures requiring treatment and measured the time from decision to treat seizures to medication administration. Shewhart charts were used for analysis.
Summary of results There were 41 seizure events among 16 patients pre-intervention (Jan 2016 to Nov 2016) and 134 events among 65 patients since the seizure rescue process was introduced (Dec 2016 to Jul 2019). Average time to seizure treatment decreased from 26 minutes when the process was not used to 12 minutes when used. Overall time to seizure treatment decreased from 25 to 13 minutes after the education initiative in Jan 2019, correlating with increased use of the seizure rescue process from 27% to 70%.

Conclusions Implementation of an innovative in-room technology was associated with a 50% reduction in time to treatment of neonatal seizures without the need to activate code resources. Education about the importance of rapid treatment of neonatal seizures increased the sense of urgency around these events resulting in more frequent use of the seizure rescue process and faster time to anti-seizure medication in the ICN.

Purpose of study Neonates may require transfer to another hospital after birth for a variety of reasons. Stabilization time (ST) is the time the transport team spends at the referring hospital. Telemedicine is an emerging technology that allows referral centers to aid in management of patients while awaiting transfer. The effect of telemedicine on ST is unknown. However, there is limited baseline data on neonatal ST for comparison. This study aims to establish baseline neonatal ST and examine how illness severity affects ST.

Methods used Retrospective chart review of transports from July 2014 to June 2017 of neonates <72 hours of age transported from a Level I/II nursery to a Level III/IV. Infants previously discharged home were not included. ST was defined as the difference between the time the transport team arrived at and departed from the referring hospital. Illness severity was measured by the Transport Risk Index of Physiologic Stability, Version II (TRIPS-II). Additional data on gestational age (GA), birth weight (BW), five-minute APGAR, and transport provider were collected to allow for statistical correction of illness severity. Statistical analysis was done using linear regression and analysis of variance (ANOVA).

Summary of results Of 668 transport charts reviewed, 472 were included in the data analysis. Mean ST was 65 ± 34 minutes (range 20-241). TRIPS-II scores ranged from 0 to 49. When TRIPS-II scores were grouped by illness severity, 385 patients fell in Group 1 (0-10), 55 in Group 2 (11-30), and 32 in Group 3 (>30). Linear regression models show statistically significant increases in ST by 1.02 minutes for every increase in TRIPS-II score by 1 point. Statistical significance was maintained when adjusting for confounders. Using grouped TRIPS-II scores, when compared to patients in Group 1, the patients in Group 2 required an extra 38 minutes of ST, and the patients in Group 3 required an extra 60 minutes of ST.

Conclusions These results indicate that ST is significantly affected by illness severity, regardless of GA, BW, APGAR or transporting provider. These are the first baseline data to establish ST. Future studies showing the impact of how telemedicine affects ST will now be possible.
Methods used This is a retrospective chart review of all pregnant women that were admitted to the Labor and Delivery Unit from the ED between June 2015 and November 2018. Patients who delivered infants or presented with threatened deliveries were included in the study. Women who had an intrauterine fetal demise were excluded. Maternal and neonatal factors were compared between the infants who survived and those that died. This study was approved by our institution’s IRB.

Summary of results A total of 71 live births occurred during the study period meeting all inclusion criteria and 7 (9.8%) of these infants died. The median maternal age was significantly younger (18 years vs. 28 years, p=0.003). All infants required resuscitation at birth. The median gestational age was 31 weeks and the mean birth weight was 2289g. The infants that died had a significantly lower gestational age (31 weeks vs. 38 weeks p=0.03). There were a total of 21 preterm infants in the study group and 5 of them did not survive. Of the infants that died, 48% had a gestational age of less than 29 weeks. Three of the preterm infants died due to failure to respond to resuscitation measures and two had congenital anomalies. There was an increased likelihood of hypoxic ischemic encephalopathy (HIE) in term infants who expired compared to the term study infants that survived (p=0.004).

Conclusions Unplanned out-of-hospital deliveries requiring resuscitation in the ED may contribute to an increased risk of mortality in preterm infants. Term infants delivered out of hospital have increased incidence of HIE with poor prognosis.

Neonatology pulmonary III
Concurrent session
10:15 AM
Saturday, January 25, 2020

470 IMPROVING SHORT TERM OUTCOMES IN VLBW INFANTS VIA LESS INVASIVE SURFACTANT ADMINISTRATION

A Diamond*, A Soliman, C Tay, S Lange, JL Kim. Miller Children’s and Women’s Hospital Long Beach, Long Beach, CA

Purpose of study Less invasive surfactant administration (LISA) has been shown to minimize the need for intubation and improve respiratory outcomes in preterm neonates by limiting exposure to positive pressure ventilation (PPV). However, the feasibility of this technique in spontaneously breathing VLBW infants and comparison with methods requiring intubation (such as INSURE) is lacking. We seek to determine whether the administration of surfactant via LISA is superior to intubation and surfactant administration by comparing short-term outcomes, and to study the applicability and feasibility of this technique.

Methods used VLBW infants meeting inclusion criteria (£34 weeks gestation, no anomaly that would impact respiration, spontaneously breathing on CPAP with evidence of RDS) received surfactant via LISA starting 04/2017. They were compared to a historical cohort receiving surfactant via endotracheal intubation. Average FiO2 requirements and level of respiratory support 12 hours post-surfactant, need for reintubation and/or additional surfactant doses, and incidence of pneumothorax within 72 hours of surfactant delivery were studied.

Summary of results Of the 117 infants in the study, 50 were in the control group (average GA 26 6/7 weeks, BW 893g) and 67 were in the LISA group (average GA 27 6/7, BW 907g). Those receiving LISA had lower FiO2 needs (0.29, 0.27, and 0.26 at 1, 2, and 12 hours post-surfactant) compared to the control group (0.35, 0.34, and 0.27, respectively). Infants receiving surfactant via LISA were less likely to require intubation 72 hours after dosing (27% vs 68% in control group).

Conclusions LISA is easily implemented, reduces average oxygen needs, and decreases the need for PPV and mechanical ventilation in VLBW infants. Given clear short-term benefits, LISA may be the preferred method for surfactant delivery in spontaneously breathing preterm infants.

Abstract 471 Table 1 CO-Hb at different DOL and BPD

<table>
<thead>
<tr>
<th>BPD n=18</th>
<th>Non-BPD n=4</th>
<th>p</th>
</tr>
</thead>
<tbody>
<tr>
<td>CO-Hb at DOL 1 to 5</td>
<td>2.47 (0.18)</td>
<td>2.85 (0.62)</td>
</tr>
<tr>
<td>CO-Hb at DOL 6 to 10</td>
<td>2.33 (0.46)</td>
<td>2.57 (0.46)</td>
</tr>
<tr>
<td>CO-Hb at DOL 11 to 14</td>
<td>2.2 (0.31)</td>
<td>2.43 (0.30)</td>
</tr>
</tbody>
</table>

Data reported as: Median (IQR)
Conclusions Lower gestational age and birth weight are associated with BPD but not CO-Hb levels at any time within the first 14 days of life. This needs to be further studied in a larger sample size.

Abstract 472 Table 1: Gain values and odds ratios

<table>
<thead>
<tr>
<th>Variable</th>
<th>Gain</th>
<th>Odds Ratio (95% CI)</th>
</tr>
</thead>
<tbody>
<tr>
<td>Head circumference at birth</td>
<td>2.37</td>
<td>0.62 (0.41 – 0.94)</td>
</tr>
<tr>
<td>Post delivery room CPAP</td>
<td>2.81</td>
<td>1.38 (1.01 – 1.89)</td>
</tr>
<tr>
<td>Days intubated ventilated</td>
<td>3.86</td>
<td>5.05 (3.01 – 8.47)</td>
</tr>
<tr>
<td>Discharge weight</td>
<td>4.04</td>
<td>1.27 (0.96 – 1.66)</td>
</tr>
<tr>
<td>Supplemental oxygen on day 28</td>
<td>14.26</td>
<td>15.29 (8.65 – 35.14)</td>
</tr>
</tbody>
</table>

Conclusions Our model found top predictors of BPD in very low birth weight infants, as well as the relative importance of these predictors. Though some of these are well-established predictors of BPD, others such as head circumference at birth and discharge weight are novel findings.
Hospital between 1/1998-6/2019. We excluded neonates repaired on ECMO.

**Summary of results** We identified 230 CDH infants repaired off ECMO. CDH neonates repaired in the NICU (n=112) were younger at birth (P<0.03), had more frequent liver up herniation (P<0.001) and less often had a primary repair (P<0.001) compared to those repaired in the OR (n=118). Age at repair was later, more often on high frequency ventilation and vasoactive medications, and had higher pre-operative oxygenation index (OI) than those repaired in the OR (all P<0.001). There was no difference in the change in OI from pre- to post-operation between the two groups. Survival was higher (97% vs 87%, P=0.007) and need for ECMO lower (2% vs 29%, P<0.001) for infants repaired in the OR. For those that died, there was no significant difference in time from operation to death between groups (table 1). By logistic regression, gestational age and defect size were significantly associated with death, while location of repair, need for ECMO and OI at 24 hours of life were not.

**Conclusions** In our center, critically ill CDH neonates are more likely to undergo operative repair in the NICU versus the OR. Cardiopulmonary function, as measured by change in pre- and post-op OI, did not differ based on repair location. Repair location was not independently associated with mortality.

### Abstract 474 Table 1

<table>
<thead>
<tr>
<th>Demographics:</th>
<th>Repair in OR</th>
<th>Repair in NICU</th>
</tr>
</thead>
<tbody>
<tr>
<td>Gestation (weeks)*</td>
<td>38±2.1±1.8</td>
<td>37±6±2.2</td>
</tr>
<tr>
<td>Liver up**</td>
<td>28±118</td>
<td>59±112</td>
</tr>
<tr>
<td>24±118</td>
<td>59±112</td>
<td>24% (53%)</td>
</tr>
<tr>
<td>Age at repair (hours)</td>
<td>58 (32-94)</td>
<td>107 (76-190)</td>
</tr>
<tr>
<td>Repair with muscle flap/patch</td>
<td>15±118</td>
<td>54±112</td>
</tr>
<tr>
<td>13±118</td>
<td>54±112</td>
<td>13% (48%)</td>
</tr>
<tr>
<td>Pre-Operative: No HF or ECMO</td>
<td>62±118</td>
<td>9±112</td>
</tr>
<tr>
<td>5±118</td>
<td>9±112</td>
<td>5% (33%)</td>
</tr>
<tr>
<td>Vasoactive medications†</td>
<td>54±116</td>
<td>78±95</td>
</tr>
<tr>
<td>54±116</td>
<td>78±95</td>
<td>54% (47%)</td>
</tr>
<tr>
<td>Oxygenation Index (OI)‡</td>
<td>4 [2.7-5.5]</td>
<td>5.7 [4.3-8.7]</td>
</tr>
<tr>
<td>Hospital Course: Change in Pre/Post-op OI§</td>
<td>-0.4 [-1.2 to 1.0]</td>
<td>-0.4 [-1.5 to 0.9]</td>
</tr>
<tr>
<td>Required ECMO</td>
<td>2±118 (2%)</td>
<td>3±112</td>
</tr>
<tr>
<td>1±118 (1%)</td>
<td>3±112</td>
<td>1% (9%)</td>
</tr>
<tr>
<td>Survived</td>
<td></td>
<td>11±112 (97%)</td>
</tr>
<tr>
<td>Operation to discharge/death (days)‌</td>
<td>22 [15-34]</td>
<td>45 [23-74]</td>
</tr>
<tr>
<td>Operation to death (days)‌</td>
<td>50 [35-89]</td>
<td>68 [21-120]</td>
</tr>
</tbody>
</table>

* Mean±SD; † n/N (%); ‡ Median (IQR 25-75%); HFV, high frequency ventilation

**Purpose of study** Prior research has shown an association between abnormal outcomes in the full term neonate due to the presence of metabolic acidemia at birth; however, few have looked into this association in premature neonates with very low birth weight (VLBW). This study aimed to describe potential association between acidemia in the premature neonate just before or at the time of delivery, with short and long term outcomes.

**Methods used** To look at the association between intrapartum acidemia and postnatal outcomes, cord blood gases as well as admission blood gases of VLBW infants were gathered between 2009 and 2019. IRB authorization was obtained from our institution prior to the start of the study. Data points were queried from NICU database as well as electronic medical records. Acidemia was defined as a pH less than 7.1 or a base excess greater than -10 mmol/L. Short and long term neonatal outcomes were analyzed against the indicator of intrapartum acidemia using SPSS statistical software Version 24.

**Summary of results** Out of 341 VLBW infants admitted to the NICU during the study period 50 infants had acidemia. Delivery room intubation and acidemia had a statistically significant relationship (p=0.005) as 65.9% of acidotic babies needed intubation compared to 42.4% of non-acidotic babies. Acidotic babies were also more likely to need chest compressions (32% vs 14% p=0.008). Severe intraventricular hemorrhage (IVH) was noted to be significantly higher (p<0.001) as 19% of acidotic babies had severe IVH compared to 3.2% of VLBW infants without acidemia. Survival rates for infants born with acidemia were significantly lower (75% vs 93%, p<0.001).

**Conclusions** VLBW infants born with intrapartum acidemia require more extensive resuscitation at the time of delivery. These infants are at higher risk of severe IVH and have increased mortality. These implications can help guide physician risk assessment of VLBW infants.

### Abstract 475

**ACIDEAMIA AT BIRTH IN THE PREMATURE NEONATE AND ASSOCIATED OUTCOMES**

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10.1136/jim-2019-WMRC.475

**Factors Affecting Need for Resuscitation in Infants of Unplanned Out of Hospital Birth and Threatened Delivery to the Emergency Room**

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10.1136/jim-2019-WMRC.476

**Purpose of study** Perinatal mortality is high in unplanned out of hospital births. The objective of this study was to analyze unplanned out of hospital deliveries and threatened deliveries presenting to the LAC+USC Medical Center Emergency Department (ED). The aim of this study was to better characterize what factors contributed to infants who needed resuscitation at birth.

**Methods used** This is a retrospective chart review of infants delivered to women that were admitted to the Labor and Delivery Unit from the ED between June 2015-November 2018 at LAC+USC Medical Center. Only patients who delivered infants outside of the hospital or presented with threatened deliveries to the ED were included. Women with an intrauterine fetal demise were excluded. This study was approved by our institution’s IRB. Data was collected for all maternal and neonatal factors through electronic medical records and an ED database.
A total of 71 live births met inclusion criteria. Of these infants, 23 (32%) of them required resuscitation following delivery. Median maternal age, positive maternal urine toxicology, and weight were not significantly different between groups. Concerning those that received resuscitation, they were more likely to have no prenatal care documented (p=0.002). The most common types of resuscitation needed included oxygen administration (74%), positive pressure ventilation (65%), and intubation (43%). Preterm infants were significantly more likely to need resuscitation compared to term infants (p=0.002). Resuscitated infants were much more likely to be admitted to the NICU than non-resuscitated infants (83% vs. 25%, p<0.001). This was still significant when correcting for term gestation (80% vs. 21%, p<0.001). In addition, resuscitated infants had a lower first temperature documented (p=0.008) and a lower initial blood glucose levels (p=0.02).

**Conclusions** Infants of unplanned out of hospital birth and threatened delivery have a higher need for resuscitation, especially if they are preterm. Lack of prenatal care increases the threatened delivery have a higher need for resuscitation, especially if they are preterm. Lack of prenatal care increases the need for resuscitation. Increased incidence of low blood glucose as well as temperature occurs in infants needing resuscitation.
A RARE CASE OF SYRINGOBULBIA IN A PATIENT WITH COCCIDIOIDOMYCOsis MENINGITis

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10.1136/jim-2019-WMRC.479

Case report Meningitis is the most extreme and lethal presentation of coccidioidomycosis and is a challenging manifestation to treat. We describe an unusual case of a young Hispanic male whose treatment for coccidioidal meningitis is further complicated by CNS anatomical abnormalities.

Methods used Retrospective case report.

Summary of results We report the case of a 20-year-old Hispanic male who was diagnosed with coccidioidal meningitis two years prior. He underwent a ventriculoperitoneal shunt procedure and endoscopic third ventriculostomy shortly after his diagnosis. After failing fluconazole treatment, isavuconazole was initiated one year later. Several months prior he experienced left upper extremity weakness and left chest, arm and face paresthesia. Evaluation by MRIs of the brain and cervical spine revealed Chiari 1 malformation with a cystic cavity in the upper cervical spine extending into the brainstem with syringomyelia and syringobulbia. The patient underwent posterior fossa craniotomy with C1 laminectomy and reported improvement of numbness and paresthesia with some residual facial numbness. The patient experienced several months without any unusual symptoms except for morning headaches which improved with over the counter medication. He recently developed shunt failure with acute diplopia and headache with progression of the syringomyelia and syringobulbia and increase in size of the dilated fourth ventricle. He underwent shunt revision with resolution of his symptoms.

Conclusions The concomitance of coccidioidal meningitis with Chiari 1 malformation, syringobulbia and syringomyelia is rare and limited literature is published. The contraindication of performing lumbar puncture to monitor treatment will make the management of this condition very challenging.
Abstracts

Here we present the case of a 41-year-old female who presented with a 8-week history of progressively worsening headache with accompanying nausea, vomiting and diplopia. She was being by ophthalmologist for several years and was diagnosed with bilateral staphyloma and myopic degeneration. Her examination showed decreased visual acuity of 20/400, bilateral cranial six nerve palsy, diffuse vitiligo patches on hand face and body and poliosis of head hair. Her lumbar punctured (LP) showed elevated intracranial pressure (ICP) of 360 mm H2O with WBC of 440, %79 lymphocytes, glucose of 56 and protein of 122. Her ICP was reduced with serial LPs. Comprehensive serum and cerebrospinal fluid studies only was positive for Quantiferon-TB Gold. She had periodic attacks of decreased visual acuity during her pregnancies. She was empirically started on four-drug standard anti tuberculosis treatment for presumptive central nervous system (CNS) toxoplasmosis. A zidovudine (AZT) MRI was performed on admission. The patient had acute encephalopathy and diffuse cerebrovascular disease involving the deep white matter, bilateral thalamus, bilateral corona radiata, and periventricular white matter with a small area of hypodensity in the left frontal subcortical white matter. This area was consistent with the ADEM. A single patient case report.

Conclusions The management of an HIV host with unknown CD4 count presenting with AMS and headaches who is found to have a CNS lesion requires immediate treatment for opportunistic infections such as toxoplasmosis. In HIV hosts with CD4 count >500, negative serum IgM/IgG for toxoplasmosis, and negative CSF DNA-PCR for toxoplasmosis, further evaluation with repeat MRI brain status post 2-3 weeks of treatment and possible brain biopsy is necessary.}

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**FRONTAL CORTEX LESION IN HUMAN IMMUNODEFICIENCY VIRUS HOST**

D Le, S Nguyen, A Abad, M Valdez*, D Aguirre. Kern Medical, Bakersfield, CA

10.1136/jim-2019-WMRC.481

**Purpose of study** In human immunodeficiency virus (HIV) hosts with CD4 cell count >500, a hyper-intense frontal cortex lesion is more likely associated with benign and malignant brain tumors than opportunistic infections, similar to findings in immunocompetent hosts.

**Methods used** Single patient case report.

**Summary of results** 57-year-old African American male with schizoaffective disorder, late latent syphilis, chronic Hepatitis C, and HIV presented with altered mental status (AMS) and headaches. The patient was diagnosed with HIV approximately 26 years prior while incarcerated. He reported that he was on and off of anti-retroviral therapy (ART) while in and out of jail and currently off ART for 7 months. CT brain with contrast was remarkable for a 1.5cm enhanced lesion with surrounding cerebral edema in the left frontal parietal lobe and subsequent MRI with contrast showed a hyperintense frontal lobe lesion. As CD4 count was unknown at this time, Bactrim was started at 5mg/kg TMP and 25mg/kg SMX for treatment of presumed central nervous system (CNS) toxoplasmosis. Azithromycin was started for Mycobacterium avium prophylaxis. Additionally, the patient was treated with Penicillin G for possible neurosyphilis. CSF analysis showed WBC 4 with 96% lymphocytes, glucose 51, and protein 32. Additional studies revealed CD4 count 565 and HIV viral load 205. Comprehensive screening in HIV-positive host, including coxi serology, toxoplasma IgG, Histoplasma urine antigen, CSF cryptococcal antigen, and CSF DNA-PCR for toxoplasmosis were found to be negative and led us to broaden the differential diagnoses to include pathologies beyond opportunistic infectious etiologies. Antibiotic therapy for treatment of presumed CNS toxoplasmosis will be continued for 2 weeks followed by repeat MRI with gadolinium. Neurosurgery will be consulted for possible biopsy if no improvement in the lesion is observed.

**482**

**INSPIRATORY STRIDOR AS AN INITIAL PRESENTATION OF ACUTE DISSEMINATED ENCEPHALOMYELOITIS**

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10.1136/jim-2019-WMRC.482

**Case report** Acute disseminated encephalomyelitis (ADEM) is a rare condition, reporting 3-4 cases per year in the USA. This case is novel because of its unique clinical presentation and rapid evolution in a pediatric patient.

The patient is a 9-year-old healthy AA female who was noted to have difficulty breathing, intermittent wheezing and inspiratory stridor 3 days prior to admission. She had two prior ER visits with normal CBC, CMP and rapid strep; a neck XR showed subglottic narrowing. Initially she was treated for croup with steroids, racemic epinephrine and albuterol and discharged home with nebulizers both times. On the day of admission, mother described the same abnormal breathing pattern as well as new onset tongue swelling and complaints of neck pain. Patient was noted to be anxious with unusual behaviors such as kicking and hair pulling. Following admission, she continued to have inspiratory stridor but was also noted to have hand flapping, moaning and a rapid change in mental status. A brain CT scan showed hypo attenuation in the left frontal subcortical white matter. Shortly after, the patient developed new onset tonic-clonic seizures and a subsequent MRI reported multiple non-enhancing white matter lesions in both frontal lobes consistent with ADEM. CSF analysis was normal. Patient was then transferred to the pediatric ICU for management.

ADEM is a demyelinating process, and overall a rare illness. Investigations suggest that the cause includes molecular mimicry of myelin autoantigens that share features with infectious organisms. 72-77% of cases have an identifiable preceding illness. The neurologic symptoms of ADEM commonly appear 4 to 13 days after an infection. This case represents a rare pediatric illness, but also is unique because of the atypical clinical presentation and rapid evolution. Physicians must maintain a wide differential for abnormal breathing patterns when patients do not respond to typical therapies, especially in the presence of anxiety-like behaviors in an otherwise previously healthy child.

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**A CASE OF PRIMARY PROGRESSIVE MULTIPLE SCLEROSIS MISDIAGNOSED AS CONVERSION DISORDER**

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10.1136/jim-2019-WMRC.483

**Case report** Psychiatric patients have their neurological conditions often overlooked secondary to underlying psychiatric disorders and are not reported well in current literature.
Methods Retrospective case report.

Results The patient is a 45-year-old Caucasian female with a history of schizoaffective disorder. The patient had a history of admissions to outside hospitals for complaints of weakness that resolved without treatment and she was diagnosed with conversion disorder. During her hospitalization, the patient reported generalized weakness, numbness, and multiple episodes of urinary and bowel incontinence. No collateral information could be ascertained to validate her symptoms. The physical examination was notable for decreased strength in her bilateral lower extremities. However, multiple hospital staff witnessed the patient moving without difficulty, making conversion disorder our diagnosis. CT brain and cervical spine revealed C3-C7 degenerative joint disease and spondylosis without any significant concern for severe neurologic impairment. A subsequent MRI of the cervical spine ruled out any concern for cervical myelopathy but MRI brain revealed periventricular lesions concerning for Multiple Sclerosis. A lumbar puncture was performed. CSF was positive for oligoclonal bands and IgG. The patient was diagnosed with Relapsing Remitting Multiple Sclerosis and started on Methylprednisolone IV 1000mg for five days. After completing Methylprednisolone, the patient made no significant improvement which prompted the initiation of plasmapheresis. The patient showed consistent improvement in her physical capacity with each session of plasmapheresis. She went from not having the ability to move from laying to sitting, to finally being able to move from laying to standing without assistance.

Conclusions Identifying Multiple Sclerosis in patients with psychiatric illnesses can be challenging, therefore as physicians it is imperative to remain diligent in identifying these features in the most complex of patients in order to detect and treat them early and properly.

Not presented – published only

484 TACKLING STIGMA AGAINST DEPRESSION AMONGST MEDICAL STUDENTS
R Kumar*, D Julian, C Ronkon, B Attridge. University of Nevada Reno School of Medicine, Reno, NV
10.1136/jim-2019-WMRC.484

Purpose of study Depression amongst medical students is prevalent due to many stressors faced during medical education. However, many do not seek treatment due to stigma from peers, self-stigma, and other barriers. A systemic review and meta-analysis by LS Rotenstein, et al (2016) suggests that 1 in 4 medical students reports being depressed. Per Schwenk et al (2010) medical students show an increase in suicidal ideation in their 3rd and 4th years, yet only 22% of depressed medical students report using mental health services. It is essential that medical students address the factors and barriers contributing to the prevalence of depression amongst medical students. This study explores methods to tackle the stigma of depression amongst medical students via interventions that aim to encourage discussion.

Methods used Peer led interventions including lectures and self-disclosure are the primary modes of intervention. Incoming medical students at UNR Med Class 2022 underwent an intervention during orientation week and during the last block of their first year. Class 2021 was used as the placebo cohort. The goal was to present the facts about depression, sources of stigma, self-awareness of mental well-being, and resources. Change in stigma was measured via a pre and post survey.

Summary of results Data is currently being analyzed for the intervention and placebo cohort. Pre-liminary results suggests an increase in the number of students diagnosed with depression in both classes. Individual questions showed a wide variability in responses. For example, the intervention group was more likely to share depressive symptoms with friends and consider medical students with depression equally as intelligent. However, they also thought they would be viewed as ‘unable to handle my responsibilities.’ Individual analysis of questions and their implications for the intervention and change in stigma is still underway. Additionally, the top 3 reasons for not seeking treatment included lack of time, stigma of being diagnosed with a mental illness, and fear of documentation on academic record.

Conclusions Due to the small sample size no statistically significant conclusion can be drawn from the data collected thus far. However, it is apparent that there was an increase in depression rates and common barriers to seeking treatment need to be addressed among medical students.

485 METABOLES IMPORTANT IN THE DIFFERENTIATION OF HUMAN INDUCED PLURIPOTENT STEM CELLS TO LUNG PROGENITOR CELLS
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10.1136/jim-2019-WMRC.485

Purpose of study Respiratory distress syndrome (RDS) affects preterm infants due to stunted lung development at birth. Induced pluripotent stem cells (iPSCs) have been successfully differentiated into lung organoids to study lung development and disease in vitro. Metabolomic technologies have allowed the examination and identification of metabolic pathways and processes occurring during development and has been successful at identifying significant biomarkers important in differentiation.

Methods used We used two wild type stem cell lines and exposed them to cocktails of growth factors and small molecules in order to undergo step wise differentiation into lung progenitor cells. The cells were sorted for CXCR4 at definive endoderm, CD271 and CD36 at anterior foregut endoderm and CPM at the lung progenitor step. The cell products were analyzed with AFBM using a Biocrates p180 kit for hexoses, amino acids, phosphatidylcholines, lysophosphatidylcholines, sphingolipids, acylcarnitines, and biogenic amines. The data was analyzed using DIABLO followed by leaps subset regression.

Summary of results DIABLO partial least squares discriminant analysis (PLS-DA) cross-compared all samples simultaneously using each differentiation state as a categorical variable. DIABLO identified signature metabolites changing between two or more differentiation states. To further refine results, top-ranked combinations of metabolites changing across pairs of differentiation stages were identified using Bayesian information criterion (BIC) scoring in subsets regression. iPSC to AFE or LPC showed unique amino acid changes (Trp, Pro) which did not appear in the iPSC to DE transition. Top-scoring unique

Conclusions These metabolites may be used in vitro to increase the efficiency of the differentiation process into lung progenitor cells and improve our understanding of early embryonic lung development. Further examination of the impact of these metabolites is warranted.

**486** IS THERE A RELATIONSHIP BETWEEN DIABETES STIGMA AND DEPRESSION?

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10.1136/jim-2019-WMRC.486

Purpose of study The relationship between diabetes and depression has been examined in several studies, with findings that suggest individuals with diabetes have higher likelihood of depression. It is important to explore this connection further, as poorer outcomes are associated with this comorbidity. One potential factor that could be involved in this relationship is diabetes stigma. Other types of health-related stigma, such as HIV-stigma, lung cancer stigma, and epilepsy stigma, have been shown to be associated with increased depression and worse health outcomes. The hypothesis is that higher diabetes stigma scores would be associated with more depressive symptoms.

Methods used This was tested in a cross-sectional survey study. Participants completed the Comprehensive Diabetes Stigma Scale (CDSS-15) to measure the experience of diabetes stigma, and the Center for Epidemiologic Studies Depression Scale-10 (CES-D-10) to measure depressive symptoms.

Summary of results A moderate positive correlation was found between diabetes stigma and depressive symptoms (r=0.41). The stigma score was significantly different between racial groups, and the depression score was significantly different between gender groups. The correlation between stigma and depressive symptoms was higher in the female group than the male group.

Conclusions Diabetes stigma could have a role in the experience of depression in patients living with Type 2 diabetes and may be a modifiable target to ultimately improve health outcomes.

**487** EXERCISE TREADMILL TESTING AND THE RATE OF FALSE POSITIVE ETTS IN WOMEN

L Mihaila*, E Amsterdam. UC Davis, Sacramento, CA

10.1136/jim-2019-WMRC.487

Purpose of study Previous studies suggest that females have a higher incidence of FP exercise ECG tests than males, thus reducing the value of this test in women. According to Sketch et al., there is significant difference in the results obtained from males and females, such that a positive ETT is of limited reliability in predicting the presence of CAD in women. However, Levisman and Amsterdam of UCD, demonstrated that the high rate of FP ETTS in women was related to the inclusion of primarily premenopausal women in whom the prevalence of CAD is low. The purpose of this study, therefore, is to investigate these divergent propositions through a retrospective review of exercise treadmill laboratory records, specifically evaluating the influence of sex and age on frequency of FP results.

Methods used Study Cohort: female and male patients with ages ranging from 30 y.o. to 62 y.o. and older, at the UC Davis Medical Center who had a positive ETT and then underwent further confirmatory cardiac testing such as exercise stress echocardiography (ESE), stress nuclear testing or coronary angiography. Also, it will include patients referred primarily for chest pain and exclude those with documented CAD, coronary angiography, myocardial infarction or revascularization.

Experimental Design: participants will be divided into 2 groups based on gender and 4 groups based on age: group I: 30-40 y.o.; group II: 41-51 y.o.; group III: 52-62 y.o.; and group IV: >62 y.o. To help identify FP ETTS the study will include parameters such as ETT-induced chest pain; age-predicted maximal HR achieved; cardiac workload; peak METs; number of ECG leads with ischemic ST depression; mm ECG ST depression; and post-ETT time to resolution of ischemic ST depression.

Summary of results I am currently collecting data and evaluating the relation of FP ETTS to the age-based subgroups and anticipate that, consistent with my hypothesis, the rate of FP ETTS will be inversely related to the age of the cohort participants.

Conclusions I am currently reviewing and analyzing study data, however some limitations we are anticipating include those inherent in the retrospective methodology as well as the collection of patient data from a single center laboratory. However, the results of this study should add to the body of knowledge on the topic of false positive ETT in women and men with potential to enhance patient management.

**488** MEDICAL MISSIONS INCREASE CULTURE COMPETENCE IN MEDICAL STUDENTS

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10.1136/jim-2019-WMRC.488

Purpose of study Culture competence is the ability to communicate across cultural differences, and is increasingly important for medical professionals. The current study aimed to assess the change of culture competence before and after a medical mission abroad.

Methods used 58 osteopathic medical students went on medical missions to Thailand, Morocco, Vietnam, or Peru between June 10 and August 7, 2019. Each participant was emailed a pre-survey 2 days pre their trip, and a post-survey 2 days post their trip to complete online using the survey platform Qualtrics. The surveys consisted of the Health Beliefs Attitude Survey questions, based on a 6-point Likert scale, where 6 stands for ‘strongly agree’ on statements of culture competence. These scores were added up for each participant, with the possible range of 15 to 90. These numbers were compared between participants pre and post survey. Wilcoxon rank sum tests were conducted to assess whether there was a
statistically significant change in the culture competence score before and after the medical mission trip. P-value<0.05 was considered to be statistically significant.

**Summary of results** 8 participants completed both pre and post survey and were included in the final analysis. More than half (62.5%, n=5) were between 26 and 30 years old, 50% (n=4) were females, 50% (n=4) were Caucasian, and 50% (n=4) attended a previous mission abroad. The median pre-score was 73.5 (Q1=67, Q3=80.5), and the median of the post score was 78.5 (Q1=74.5, Q3=82.5). There is a marginally statistically significant change in the median between pre and post survey (p=0.0938). Among participants who attended a medical mission before this study, the culture competency score increased from 78.5 to 80.5 (delta=2, p-value=0.3750). Among participants who did not attend a medical mission before, the culture competency score increased from 69 to 75.5 (delta=6.5, p-value=0.5000).

**Conclusions** Due to small sample size, the change in the median of culture competency score did not reach statistical significance. However, the trend of increased culture competence demonstrated the effectiveness of a medical mission abroad, especially for those who did not attend a medical mission abroad in the past. Future research based on a larger sample size should be conducted to confirm the current findings.

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**APPENDICITIS (NOT!): CECAL DIVERTICULITIS IN AN ADOLESCENT MALE**

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**Introduction** While diverticular disease is typically thought to be a disease of the elderly, recent epidemiological studies demonstrate increasing incidence among patients younger than 50 years old. Even so, diverticulitis is still exceedingly rare within the pediatric population.

**Case report** We present the case of a 15 year old previously healthy male who presented with four days of worsening sharp right-sided mid-abdominal pain associated with nausea, constipation, and poor appetite. His labs were notable for an elevated C-reactive protein and mildly elevated ESR but otherwise normal CBC and CMP. Pediatric Surgery was consulted due to concern for appendicitis. He underwent an abdominal CT which demonstrated a single cecal diverticulum containing high attenuation material with thickening of the cecal wall and associated fat stranding. While the appendix seemed somewhat enlarged, it was partially gas-filled. His clinical picture and imaging were consistent with diverticulitis. He was started on ceftriaxone and metronidazole. Diet was slowly advanced as tolerated. His pain improved over the course of his 4-day admission. Aside from one temperature to 38 degrees F, he remained afebrile throughout hospitalization. He was discharged home with amoxicillin/clavulanic acid in order to complete a 7-day antibiotic course.

**Conclusion** Diverticulitis should be considered as a potential diagnosis when pediatric patients present with acute onset of severe abdominal pain, especially in presentations that are concerning for but may not fully fit an appendicitis illness script.

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**A CASE OF GLIOMATOSIS CEREBRI**

N Hasan*, D Aguirre, S El-Halees. Kern Medical, Bakersfield, CA

**Introduction** Glioblastoma cerebri is an extremely rare and aggressive brain tumor with resistance to therapy. Incidence reported to be less than 100 cases each year in the USA.

**Discussion** 49 yo male with a history of IV meth abuse, ETOH abuse, and seizure disorder on Keppra 750mg twice daily presented with increasing frequency of seizures. He reported having five episodes daily involving upper extremity clenching with loss of consciousness and urinary incontinence. Seizures were typically followed by 4-5 minute post ictal period. He also reported daily visual hallucinations for the past 3 weeks. Medical records from 1 month prior indicated an unremarkable CT and MRI brain, no epileptiform activity on EEG monitoring and negative CSF analysis for infections. Labs including toxilology were unremarkable on presentation. Overnight EEG monitoring indicated no epileptic activity. MRI brain was notable for multiple areas of subtle white and gray matter T2/Flair abnormalities in the bilateral frontotemporal lobes, with one discrete small 7 mm enhancing ring lesions in the left temporal area, highly suspicious of gliomatosis cerebri type 2.

Neurosurgery was consulted for brain biopsy and pathology of the right temporal lobe was positive for diffuse astrocytoma (WHO grade II/IV). Patient was started on Depakote for seizures as well as chemotherapy and radiation for four months. Post radiation, MRI brain was repeated and showed worsening bilateral frontotemporal hyperintensities, and an enlarging left temporal lesion with three new left frontal lesions. Oncology referred the patient to neurosurgery for consideration of resection, however were unfortunately notified that patient had already deceased.

**Conclusion** Gliomatosis cerebri involves multiple contiguous lobes of the brain. Main symptoms are dementia, seizures, and hemiparesis. Diagnosis is made with MRI and histopathologic confirmation of an astrocytic process. Prognosis is poor despite aggressive treatment with 26-52% surviving less than 1 year. Surgical resection is generally not an option due to multilobar involvement, radiation and chemotherapy has no proven efficacy. Major obstacles toward establishing standardized treatments include rarity of the disease, lack of understanding of the tumor biology, variation across histopathologic grading and patient outcomes, and lack of response to therapies.

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**A CASE OF HGB S/O ARAB’S DISEASE IN A HISPANIC MALE, CASE REPORT AND LITERATURE REVIEW**

BA Mocan*, T Bajaj, E Davtyan, J Patel, A Prunes. Kern Medical-UCLA, Bakersfield, CA

**Case Report** Hemoglobin S/O Arab’s disease is seen primarily in those of Middle Eastern descent. To the best of our knowledge, there is no reported or published case of this disease in the Hispanic population.
A 26 year old male diagnosed at 2 months of age with Hemoglobin S/OArab presented to the emergency department with severe abdominal and right leg pain. Several days prior to hospitalization, he had intractable nausea and non-bilious, non-bloody vomiting. His abdominal pain worsened and spread to his bilateral hips and proximal thighs, prompting him to seek medical attention. He had stopped his daily hydroxyurea and folic acid approximately 3 months prior. On physical exam, he was in acute distress with scleral icterus and epigastric tenderness reproducible on palpation. A chest x-ray illustrated no acute cardiopulmonary disease. Laboratory results were significant for leukocytosis, normocytic anemia, reticulocyte count of 9.89%, absolute reticulocyte count of 5.9, total bilirubin of 7.1 (conjugated 1.4), and an LDH of 962. Peripheral blood smear showed hypochromasia, polychromasia, sickle cells, and target cells. The patient’s pain was controlled with hydrocodone 5mg/acetaminophen 325mg and 2 mg intravenous morphine for breakthrough pain control. He was restarted on folic acid 1 mg daily and hydroxyurea 500mg twice daily. Pain was adequately controlled and there was no evidence of underlying infection. Patient was discharged with the same regimen and followed closely in outpatient clinic.

Hemoglobin S/OArab is a rare hemoglobinopathy caused by two variant beta globin chains with glutamate replaced by valine and lysine. The presence of these two substitutions results in a presentation comparable to sickle cell disease. Hb S/OArab vaso-occlusive crises can be managed similarly to those of Hb S. However in our case, the resolution occurred within 48 hours, which is significantly sooner than those with typical hemoglobin S disease.

We present the first documented and reported case of Hb S/OArab in a Hispanic male. The gene frequency of Hb S/OArab is unknown, though, per Zimmerman et al., of the estimated 1500-2000 sickle cell infants born each year, it is possible that several of them are Hb S/OArab cases.
Conclusions Thorough investigation should always be done for any patient who presents with neutropenic fever after chemotherapy. Most clinicians often jump to infectious causes such as opportunistic pathogens, however structural defects may be the cause of underlying illness. Due to his palatal defect caused by erosion from the lymphoma, this patient required surgical intervention as well as antibiotics to treat his complications.

**VIBRIO VULNIFICUS: WHEN OYSTERS AREN’T APHRODISIACS**

1AC Bakshi*, 2S Bakshi, 7C Cange. 7Western University of Health Sciences, Pomona, CA; 2St. George University, West Indies, Grenada

10.1136/jim-2019-WMRC.494

**Case report Vibrio vulnificus** is a gram negative bacteria can potentially cause serious infections and septicemia. With approximately 200 reported infections each year, it is an easily overlooked diagnosis. Delays between onset of symptoms and initiation of treatment have been associated with higher mortality rates which makes early identification crucial.

A 43-year-old male presented to the emergency department at St. Francis Medical Center, Lynwood, CA with a 4-day history of weakness, nausea, and bilateral lower extremity pain. The patient denied any medical history but admitted alcohol abuse. Vitals consisted of blood pressure of 100/72 mmHg, heart rate of 118 bpm, respirations of 32/rpm, and a temperature of 100.4F. Physical exam showed scleral icterus, yellowing of the tongue, and bilateral lower extremities edema with erythematous macular blanching. Laboratory results included sodium of 120 meq/L, potassium of 5.6 meq/L, WBC count of 1.2 x 10^9/L, platelets of 80 x 10^9/L, BUN of 55 mg/dL, creatinine of 6.2 mg/dL, glucose of 92 mg/dL, alkaline phosphatase of 210 U/L, albumin of 3 g/dL, and total bilirubin of 1.4mg/dL. Imaging revealed hepatosplenomegaly and fatty liver. Diagnosis of sepsis was made; hypotension persisted despite vasopressors. Blood culture revealed *Vibrio vulnificus*. Doxycycline and ceftazidime was administered. Upon questioning, patient admitted to eating raw oysters in Mexico recently. Despite resuscitative measures.

The patient deteriorated over 39 days, enduring septic shock, respiratory and kidney failure, hepatic encephalopathy, and necrotizing fasciitis. He expired following a myocardial infarction despite resuscitative measures.

Although the incidence of *V. vulnificus* is low, the scenario of septicemia and bullous skin lesions should raise suspicion in patients with underlying liver disease. Susceptibility is due to higher iron levels, providing thriving environment for *V. vulnificus*. Questioning of raw seafood consumption should be considered in septic patients with cellulitis. This case illustrates the importance of obtaining a complete history and identifying the organism early. Recognition and aggressive use of supportive measures are critical in minimizing the fatal consequences of *V. vulnificus*.

**COLLOID MILIUM: A NOVEL CASE FOLLOWING THERMAL BURN WITH DIFFERENTIAL DIAGNOSIS**

1A Savell*, 2M Heaphy. 1University of Nevada, Reno School of Medicine, Reno, NV; 2Skin Cancer and Dermatology Institute, Reno, NV

10.1136/jim-2019-WMRC.495

**Introduction** Colloid milium is a rare condition of colloid deposition which most commonly affects fair skinned individuals with history of long-term sun exposure. There are no reports of colloid milium following thermal burn in the literature. Differential diagnosis of colloid milium may include conditions such as sebaceous hyperplasia, nodular amyloidosis, and basal cell carcinoma.

**Case Presentation** A 63 year-old man presented to the dermatology clinic with a papular lesion on the bridge of his nose. The papules appeared over 20 years earlier following a severe thermal burn and have been gradually improving for the last 10 years. However, his symptoms of burning sensations flare every year when he is exposed to heat and sun. He has applied high SPF sunscreen without relief. He has no known allergies. Shave biopsy was performed, and hematoxylin and eosin staining revealed homogenous, acellular material consistent with colloid extending through the dermis and exhibiting characteristic fissure artifacts.

<table>
<thead>
<tr>
<th>Abstract 495 Table 1</th>
<th>Selected differential diagnosis of colloid milium</th>
</tr>
</thead>
<tbody>
<tr>
<td>Condition</td>
<td>Characteristics</td>
</tr>
<tr>
<td>Colloid Milium</td>
<td>Deposition of colloid in dermis; small flesh colored or translucent papules most commonly on sun exposed areas of head and neck</td>
</tr>
<tr>
<td>Nodular amyloidosis</td>
<td>Cutaneous manifestation of sarcoidosis; waxy papules</td>
</tr>
<tr>
<td>Sebaceous</td>
<td>Benign hair follicle tumor; small yellow papules seen in older adults</td>
</tr>
<tr>
<td>Hyperplasia</td>
<td>Typically on forehead or cheeks</td>
</tr>
<tr>
<td>Basal Cell</td>
<td>Neoplasm of basal cells of epidermis; slow growing flesh-colored, pink, or pigmented plaque or nodule that may spontaneously bleed or ulcerate</td>
</tr>
</tbody>
</table>

**Implications** Here we present a novel case of colloid milium following a thermal burn. We outline the clinical and histopathologic differences among colloid milium and other conditions commonly included on a differential diagnosis.

**Abstract 495 Figure 1** Colloid milium on nose
Purpose of study

Newborns being evaluated in the emergency room has dramatically increased over the past several years. Neonatal susceptibility to infections, parental anxiety, insufficient caretaker knowledge, or benign symptoms are just a few reasons for the increase in the number of emergency department (ED) visits. However, several studies show that majority of newborns’ ED visits were considered non-urgent. To prevent non-urgent ED visits, it is vital that outpatient physicians provide excellent anticipatory guidance to parents of newborns. The objective of this retrospective study is to determine factors associated with newborn ED visits at Community Regional Medical Center (CRMC) and improve content, delivery, and emphasis when providing anticipatory guidance.

Methods used

We collected data of over 500 newborn seen in the ED of CRMC in Fresno, California between January 2014 and December 2018. Participants were families of term newborns ages 1-28 days, with no history of prematurity, prolonged NICU stay, or complex medical history. As a part of this retrospective study, SPSS analysis will be conducted on the following data: demographics of the newborn (gestational age, age of visit to ED) and maternal data (maternal age, number of live births, length of stay postpartum, home postal code at time of ED visit, ethnicity, race). The data will also juxtapose the parent’s chief complaint in conjunction with the physician’s final diagnosis to determine the urgency of the visit.

Summary of results

The data will identify the demographics of mothers and newborns seen in the ED, particularly mothers that are primiparous, teenagers, living in low socioeconomic areas, or those discharged early from postpartum. When categorizing parents’ main concern based on organ system and chief complaint, this breakdown will affect how we introduce a tool for providers to improve outpatient patient education.

Conclusions

By profiling the type of newborns’ visits to the ED, physicians can better understand parental concerns during the newborn period and how to provide anticipatory guidance in the outpatient setting. Furthermore, the knowledge gained from this study will help physicians improve newborn parent education in the postpartum setting to reduce the number of non-acute ED visits.

Abstract 497

INITIAL EMPRIRICAL ANTIBIOTIC EXPOSURE IN PREMATURE INFANTS AND THE RISK FOR NECROTIZING ENTEROCOLITIS (NEC)

Purpose of study

Empiric antibiotics are commonly used on premature infants during the early postnatal period. Alteration of bacterial flora in the neonate’s gut may be a risk factor for the development of NEC. The purpose of this study was to investigate whether initial empirical antibiotic treatment in premature infants is associated with an increased risk of NEC.

Methods used

A literature review was conducted using PubMed, Google Scholar, and reference checks. Keywords used included, ‘necrotizing enterocolitis,’ ‘antibiotic exposure,’ and ‘prematurity.’ Only studies that included a control group and accounted for potential confounding variables associated with NEC were included in our analysis.

Abstract 497 Table 1

Association of IEAT and development of NEC

<table>
<thead>
<tr>
<th>First Author &amp; Year</th>
<th>Mean Gestational Age (weeks)</th>
<th>Definition of Cases &amp; N</th>
<th>Definition of Controls &amp; N</th>
<th>Outcome</th>
</tr>
</thead>
<tbody>
<tr>
<td>Raba, 2019</td>
<td>NEC: 25.2 Control: 25.9</td>
<td>Infants exposed to IEAT w/NCE: 22</td>
<td>Matched controls exposed to IEAT w/o NCE: 32</td>
<td>PIEAT was associated with a 3.6-fold increase in the risk of a baby developing NEC (Odds Ratio: 3.6; 95% CI 1.13-11.47)</td>
</tr>
<tr>
<td>Esmaeilizand, 2017</td>
<td>NEC: 25.9 Control: 25.9</td>
<td>Infants exposed to IEAT w/NCE: 224</td>
<td>Matched controls exposed to IEAT w/o NCE: 447</td>
<td>Meropenem and gentamicin with highest association with NEC</td>
</tr>
<tr>
<td>Alexander, 2011</td>
<td>NEC: 28.2 Control: 28.1</td>
<td>Infants exposed to IEAT w/NCE: 124</td>
<td>Matched controls exposed to IEAT w/o NCE: 248</td>
<td>PIEAT was associated with increased risk of NEC compared with IEAT (adjusted OR: 2.02; 95% CI 1.55, 3.13)</td>
</tr>
<tr>
<td>Cotten, 2009</td>
<td>PIEAT: 25.6 Control: 26.3</td>
<td>Infants exposed to PIEAT w/NCE: 255</td>
<td>Infants exposed to PIEAT w/o NCE: 1892</td>
<td>Odds ratios of NEC with duration of IEAT: No IEAT: 1.39 [3-4 days: 1.52</td>
</tr>
<tr>
<td>Abdel Ghany, 2012</td>
<td>PIEAT: 30.9 Control: 32.6</td>
<td>Infants exposed to PIEAT w/NCE: 34</td>
<td>Infants exposed to PIEAT w/o NCE: 173</td>
<td>Antibiotic exposure was a risk factor for NEC, p=0.015</td>
</tr>
<tr>
<td>Torres, 2018</td>
<td>PIEAT: 27.8 Control: 28.2</td>
<td>Infants exposed to PIEAT w/NCE: 130</td>
<td>Infants exposed to PIEAT w/o NCE: 69</td>
<td>The duration of IEAT and risk of NEC, OR: 1.31, p=0.018</td>
</tr>
<tr>
<td>Kuppa, 2011</td>
<td>PIEAT: 27.4 Control: 29.3</td>
<td>Infants exposed to PIEAT w/NCE: 130</td>
<td>Infants exposed to PIEAT w/o NCE: 69</td>
<td>PIEAT in 34 (100%) of those with NEC vs 139 (80%), p=0.005</td>
</tr>
<tr>
<td></td>
<td></td>
<td></td>
<td></td>
<td>20 (13.9%) got NEC, all belonged to the PIEAT group, p=0.03</td>
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<tr>
<td></td>
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<td></td>
<td></td>
<td>8 (4.6%) of those in IEAT vs 9 (6.9%) of those in PIEAT got NEC, p=0.11</td>
</tr>
</tbody>
</table>
A CASE OF CONGENITAL CENTRAL HYPOVENTILATION SYNDROME COMPLICATED BY INTRACRANIAL HEMORRHAGE

M Romboy*, 1G Schaub, 2D Reyes. 1University of Las Vegas, Nevada, Henderson, NV; 2University Medical Center, Las Vegas, NV

Background Congenital Central Hypoventilation Syndrome (CCHS) is a rare cause of apnea in the neonate. We present a unique case of CCHS to demonstrate that a high index of suspicion for CCHS is necessary to avoid a delay in diagnosis.

Case report A full term baby boy was born to a healthy 25-year-old G1P1 via caesarean section with vacuum-assist due to failed induction. Patient required positive pressure ventilation at birth due to poor respiratory effort. At 3 hours of life, patient had a cyanotic episode and was admitted to the NICU. He required intubation due to apnea. Due to history of traumatic delivery, head CT was done and demonstrated bilateral subdural hemorrhages and a non-displaced skull fracture. MRI showed posterior subdural bleed. Neurology and neurosurgery were consulted to help explain the frequent apneic episodes. Patient was extubated and maintained on nasal non-invasive ventilation but continued to have frequent apnea. He failed a trial of caffeine and required reintubation due to respiratory acidosis. Subsequently, a sleep study was done revealing multiple episodes of central apneas and hypopneas. Genetic testing for CCHS was ordered and was positive for a mutation in the PHOX2B gene. Patient underwent tracheostomy and was discharged on a home ventilator.

Discussion CCHS is a rare disorder of respiratory and autonomic dysregulation estimated to affect 1:200,000 people. It is classically characterized by hypercapnia and hypoxemia during sleep. Most patients require tracheostomy and lifelong ventilatory support. Apnea in the neonate has a broad differential including cardiac, pulmonary, infectious and neurogenic causes. CCHS is often overlooked because of overlap with more common disease processes and low incidence of the disease. Our patient had a neurological explanation for apnea (head trauma with intracranial hemorrhage) and thus CCHS was not initially considered as a diagnosis. Only when our patient was unable to be extubated due to persistent respiratory acidosis was genetic testing done. Due to the availability of genetic testing for CCHS, we recommend considering CCHS as a diagnosis in all term neonates with central apnea despite presence of confounding findings.
Abstracts

500 PATIENT PREFERENCES IN COLORECTAL SURGERY: WEIGHING SURGICAL SITE INFECTIONS AND ANTIBIOTICS SIDE EFFECTS

FS Brecha*, J Esplin, E Ozanne, J Cohan. University of Utah, Salt Lake City, UT

Purpose of study The likelihood of a Surgical Site Infection (SSI) after colorectal surgery is reduced when using a mechanical bowel preparation with oral antibiotics (MBP + ABX), ABX may cause side effects such as vomiting; thus, this study aimed to understand the willingness of patients to take ABX despite the possibility of side effects to reduce the risk of SSI.

Methods used This was a single-center study of adults undergoing colorectal surgery, colonoscopy, or treatment in a surgical clinic. A standard gamble instrument was designed to measure patient willingness to accept ABX side effects to reduce the risk of SSI from 14% to 7%. Data were analyzed using the Kruskal-Wallis and Wilcoxon signed-rank tests due to non-parametric distribution.

Summary of results We enrolled 68 patients (36 women) undergoing colonoscopy (n=43), colorectal surgery (n=14), and evaluation in clinic (n=11). Willingness to take ABX decreased with increasing severity side effects, p<0.001 (figure 1). The highest level of severity of side effects patients were willing to accept was moderate (55.9%), followed by severe (23.5%), mild (14.7%), none (1.5%). Five patients that volunteered that they had personal experience with SSI chose to always complete antibiotics regardless of side effects.

Abstract 500 Figure 1 Patient willingness to take ABX assuming varying severity of side effects. Answers varied significantly using Kruskal-Wallis test, p<0.001.

Conclusions Increasing side effect severity is associated with decreased patient willingness to take ABX even with reduced risk of SSI. This effect may be modified in patients who previously experienced SSI. Further studies are needed to inform preoperative discussions about the quality of life associated with both ABX and SSI prior to colorectal surgery.

501 OPTIMIZING SKIN ANTISEPSIS FOR NEONATAL SURGERY

S Carr*, C Gogal, E Skarsgard. 1University of British Columbia, Vancouver, BC, Canada; 2BC Children’s Hospital, Vancouver, BC, Canada

Purpose of study Preoperative skin antisepsis is routinely performed to reduce the risk of surgical site infections (SSIs). Povidone-iodine has been the antiseptic of choice for decades; however, a growing body of evidence in adults suggests that alcohol-based chlorhexidine solutions provide superior protection against SSIs. Equivalent data in the neonatal surgical population is lacking, but literature on skin antisepsis for invasive non-surgical procedures in neonatal intensive care units (NICUs) and for prevention of newborn nosocomial infections in developing countries suggests that chlorhexidine antiseptics is safe and non-inferior to povidone-iodine in reducing hospital acquired infections.

The aim of this project is to therefore assess the safety (avoidance of skin injury) and effectiveness (avoidance of SSIs) of preoperative skin antisepsis with chlorhexidine-alcohol in neonates undergoing abdominal and thoracic surgery at British Columbia Children’s Hospital (BCCH).

Methods used Our target population will include general surgery patients in the NICU at British Columbia Women’s Hospital (BCWH) older than 34 weeks post conceptual age. Patients with known allergies to chlorhexidine gluconate (CHG) or isopropyl alcohol (IPA), open wounds, or mucosal surfaces in the prep area will be excluded. Patients will be prepared with one of two CHG-IPA formulations (2% CHG – 70% IPA or 0.5% CHG – 70% IPA) and skin integrity assessments will be performed pre- and post-application. Using BCCH data from the American College of Surgeons National Surgical Quality Improvement Pediatric Program, an analysis will be performed comparing pre- and post-implementation data to assess incidence of adverse skin reactions and SSIs.

Summary of results Implementation of the proposed CHG-IPA formulations is ongoing. Preliminary results will be reported at the time of the conference.

Conclusions This project will: 1) engage stakeholders from BCCH and BCWH in neonatal surgical quality improvement work, and 2) inform the development and implementation of a neonatal surgical skin preparation protocol that can be applied across surgical subspecialties to standardize practice and improve clinical outcomes (specifically reduce SSIs).

502 A RARE CASE OF PNEUMATOSIS CYSTOIDES COLI PRODUCING ILEOCOLIC INTUSSUSCEPTION

M Haseeb*, R Krishnankutty, N Yazzie. 1University of Arizona College of Medicine–Tucson, Tucson, AZ; 2Tehaatsaal Medical Center, Fort Defiance, AZ

Introduction Ileocolic intussusception is primarily a disorder of early childhood, with 10% of cases occurring in children over 5 years of age. In this older age group, it is associated with a pathologic lead point, such as lymphoid hyperplasia or a tumor. Pneumatosis cystoides intestinalis (PCI), a rare condition in which gaseous cysts line the intestinal wall, can seldom cause intussusception. Here, we present a unique case of intussusception secondary to PCI in a 7-year-old.

Case An otherwise healthy 7-year-old Navajo boy with a recent episode of self-limited constipation and abdominal pain presented to the ED with a six-hour history of colicky, periumbilical pain and diarrhea. On presentation, the patient was in acute distress secondary to pain. Contrast CT showed an
intussusception beginning proximal to the terminal ileum, extending at least 7.3 cm distally. Coronal images showed terminal ileum, cecum, and ascending colon invaginating into the distal ascending and transverse colon. Free fluid was seen at the origin of the intussusception, and general surgery was consulted for immediate reduction by exploratory laparoscopy. During surgery, the intussusception was reduced by lifting the transverse colon and allowing gravity to pull the ileum out of the ascending colon. A firm mass in the cecal wall was identified as a possible lead point. The laparoscopy was converted to an open ileocecectomy due to concern of malignant pathology. Gross pathologic examination of the cecum showed areas of dilatation with multiple air-filled cystic cavities with smooth internal linings and no mural nodularities. Histologic examination of the tissue sections was appropriately reactive. PCI was determined to be the cause of intussusception.

Discussion The association between PCI and intussusception is extremely rare, with only six documented cases in the pediatric population to our knowledge. Of these, our patient represents the second pediatric patient without an underlying secondary pathology. Despite its rarity, PCI should be considered when evaluating patients for intussusception—prompt recognition can aid in a more conservative management of the disease.

Purpose of study Women have been underrepresented in orthopedic surgery for many years despite increasing proportions of graduating female medical students and percentage of women in other surgical specialties in the recent years. Studies, however, reveal this disparity is due more to women’s decision not to pursue careers in orthopedic surgery rather than their inability to match into training programs. A mentor was defined as someone who provides guidance, advice and support in professional and personal development. Our study examined importance of same-sex mentorship at all stages of medical training, from medical students to practicing physicians.

Methods used Between May-October of 2019, UC Davis medical students, residents and faculty completed an online Qualtrics survey asking about their background, mentorship experience, and important factors when choosing a medical specialty.

Summary of results Preliminary results show responses from 324 medical students, 309 residents, and 157 faculty members. Initial data revealed that respondents generally agreed that having a mentor who takes the time to get to know their mentee matters more than having a mentor who matches their mentee’s gender. Students expressed their desire for mentors be aware of the challenges that gender might play while in pursuit of their goals. These findings support the notion that mentors can be successful even when they are not the same gender as their mentee. Most respondents believed that mentors are important, with residents valuing mentorship more now compared to earlier in their training. Several students reported that they wished they had mentors in earlier years of medical school, suggesting that medical schools should adopt programs to pair students with mentors as early as possible. Faculty members requested more protected time to spend with their mentees to build relationships and accomplish goals together.

Conclusions There is a clear demand for mentoring programs to direct focus on improving the quality of mentorship by diversifying pool of mentors and connecting mentees to the appropriate mentors they can identify with, as well as, providing protected time for mentors to cultivate the most effective relationships and success in their careers.

Abstract 504 Figure 1 Showing the 3 x 3 cm herniation of the left lobe of the liver overlying the left chest wall
Computer Tomography (CT) scans suggested the presence of a left diaphragmatic hernia, left rib agenesis and herniation of an unidentifiable intra-abdominal organ through the anterior left chest wall. On day of life (DOL) 30, when global health outreach pediatric surgeons arrived to the hospital, the decision was made to operate on the child. The mass was found to be of liver origin, the exposed tissue was excised and primary closure of the chest wall was accomplished. The patient’s postoperative course involved a wound infection that resolved favorably with treatment, allowing for discharged home on post-operative day (POD) 17 in a stable condition.

505 IMPACT OF FUNCTIONAL TOTAL LARYNGECTOMY ON QUALITY OF LIFE

JC Mecham*, SR Hall, DG Lott. Mayo Clinic Scottsdale, AZ; Mayo Clinic Arizona, Phoenix, AZ

10.1136/jim-2019-WMRC.505

Purpose of study It is estimated that 5-11% of total laryngectomies performed in the United States are done for organ dysfunction, termed functional total laryngectomy (FTL). Despite this, little has been written about the effect this has on quality of life (QOL). In this study, we aim to evaluate the impact on quality of life of this major elective surgery with life-altering effects.

Methods used A cross-sectional analysis of FTL patients was performed using the MD Anderson Dysphagia Index (MDADI), University of Washington Quality of Life Index (UW-QOL), and the 12-Item Short Form Healthy Survey (SF-12) version 2. Patients were asked to fill out the survey based on their preoperative and postoperative symptoms. Finally, they were asked if it was worth it.

Summary of results A total of 8 out of 15 patients met inclusion criteria, were able to be contacted, and were willing to participate in the study. There was no significant difference between global preoperative and postoperative QOL scores as measured by any of the three questionnaires used. When broken down by individual questions, there was a significant decrease in appearance scores (p=0.03) and a significant increase in swallow scores after FTL as measured by the UW-QOL (p=0.008). Six of the eight patients stated that they would elect to under FTL again.

Conclusions Functional total laryngectomy appears to have a neutral effect on QOL with some areas improving and others worsening. This is important information to discuss when counseling patients with end-stage dysphagia.

506 SURGICAL INTERVENTIONS FOR MITRAL VALVE BACTERIAL ENDOCARDITIS IN PEDIATRIC PATIENTS: A SYSTEMATIC REVIEW

D Rodriguez*, C Rosenow, M Frey, IJ Jahanyar. Mayo Clinic Alix School of Medicine, Scottsdale, AZ; Midwest University Chicago College of Osteopathic Medicine, Glendale, AZ; Mayo Clinic Phoenix, AZ

10.1136/jim-2019-WMRC.506

Purpose of study Bacterial endocarditis is an infection of the heart’s endothelial lining, causing the formation of bacterial vegetations most commonly around the lining of the mitral valve. In 52% of cases, surgical intervention is required. Mitral valve (MV) repair and replacement are the most common corrective procedures, however, there is scarce literature comparing the outcomes of both procedures in a pediatric cohort.

Methods used A systematic literature review using PubMed, Ovid EMBASE, Cochrane Database of Systematic Reviews, Scopus, and Web of Science identified 20 case reports for data extraction. Papers were selected through the systematic use of search terms and were manually assessed for relevance and inclusion of pertinent data. Data parameters included postoperative complications and outcomes. Review and meta-analysis were conducted according to PRISMA guidelines.

Summary of results The mean age of the patients was 6.4 years. 10 patients underwent MV repair and 10 underwent MV replacement. 1 patient had 2 repairs of the MV that resulted in the failure of each procedure. This ultimately led to the replacement of the previously repaired valve. Although MV repair had more overall complications, MV replacement resulted in more severe complications such as heart block (10%) and valve thrombosis (40%). In the MV repair group, MV regurgitation and MV stenosis were present post-operatively in 40% and 20% of patients, respectively.

Conclusions We recommend that MV repair should be the treatment of choice since complications are less grave, as they are mainly comprised of residual regurgitation or novel stenosis. The major adverse outcomes seen in patients after valve replacement are associated with more severe sequelae and future implications for patients. Improving outcomes for MV repair in this patient cohort should be the focus of possible future studies.
Correction: 2020 Western Medical Research Conference

Since initial publication of these abstracts there are some changes and additions required as follows:

doi: 10.1136/jim-2019-WMRC.135
The author order was listed incorrectly with S Ravaie as the third author. S Ravaei should have been listed as the first author. The author list should therefore be S Ravaei, K Ludwig, H Kornblum.

doi: 10.1136/jim-2019-WMRC.464
This abstract has been retracted. The online version of the abstract supplement and author index have been updated to reflect the changes.

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